

BAW-4

PICATINNY ARSENAL
TECHNICAL REPORT NO. 2510

DICTIONARY OF EXPLOSIVES, AMMUNITION AND WEAPONS

(GERMAN SECTION)

BASIL T. FEDOROFF

HENRY A. AARONSON

GEORGE D. CLIFT

EARL F. REESE

DOVER, NEW JERSEY

1958

TABLE OF CONTENTS

	PAGES
Foreword	1-IV
List of German Explosives, Ammunition and Weapons	1-264
Vocabulary of German Ordnance, etc.	285-308
List of German Abbreviations of Ordnance and Related Terms	309-346

LIST OF GERMAN EXPLOSIVES, AMMUNITION AND RELATED ITEMS

"121" (Firing Composition). See Firing Composition 121.

"A" (Rohstoff). "A" (Rohstoff). Beginning about 1933, the Germans started to experiment with military rockets. The first model was the A-1 which weighed about 330 lbs and was 17 1/2" long and 1 1/2" in diameter; it was unsuccessful. The next rocket, the A-2, which appeared in 1934, was an improved A-1 and when fired it attained an altitude of 6000 feet. In 1938, at Peenemünde, the A-3 was developed. This was the predecessor of the A-4, developed in December, 1940 and was commonly known as the V-2. The A-3 rocket weighed 1,600 lbs and was 25 1/2" long and 2 1/2" in diameter. The A-4 rocket is briefly described under V-2. The next A rocket was developed at Peenemünde; the A-5, A-6, A-7, A-8, A-9 and A-10, were purely experimental. Among them, the A-5 and A-10 were intended for bombardment of the U.S.A. The A-5 was intended to be carried aloft by the A-1 during the first phase of the near-Atlantic trip.

Reference: F. Kuntz, "Guided Missiles, Rockets and Torpedoes," London & Co., N.Y. (1951), pp. 22-34.

A-2 Same as V-2.

A-3 (Rohstoff). Same as V-2 (Rohstoff).
[See also V. Dornberger, W. S. Viking, N.Y. (1950).]

"AG" (Fuselage) were two-variant fuselages developed at Technische Fabrik an subsidiaries for the "G" 1 fuselage after it became difficult to obtain corium - magnesium alloy (called Magnesium) one of the essential ingredients of G 3.

The AG fuselages were made by dipping the tip of a bomb or rocket into the following composition:

- (1) First dip composition consisted of dry Pb picrate 50g and silicon (particle size 20 to 40 microns) 10g, suspended in about 75 ml of a 28 soln of MC in naph or hept. solvent. After dipping was dry, the head on the bridge wire was dipped into the MC.
- (2) Second dip composition which consisted of dry Pb picrate 50g, Pb chromate 35g and silicon (size 20 to 40 microns) 10g, suspended in about 75 ml of 28 soln of MC in naph or hept. solvent. The dried head was dipped into the MC.
- (3) Third dip composition which was a lacquer consisting of a 15% soln of MC in 75/75 - hept. acetone/alcohol, to which was added 20% of a 28 soln of MC in naph or hept. solvent. The dried head was dipped into the MC. The two dried heads were dipped into the MC.
- (4) Fourth dip composition which consisted of MC lacquer in (c) to which was added 0.5g of Sudan Brown (0.5g per 10 c) of lacquer.

Further operations are the same as described under Fused Glass Manufacture.

Reference: R. Ascher, et al. (OS Film Rept No. B55, Nov. No. 2 (1946), p. 4-17).

A-3/A-10: Long range guided missile designed to have a range of 7,000 km is briefly described in TM 9-1985-2 (1953), p. 233.

Abbreviations for Ordnance Terms. See Ordnance Terms and Abbreviations in this section.

Abbreviations or Abkürzungen. See Wasse (or Spent) Acids.

Abnehm 2. Same as Filler No. 57.

Absolute Method of Measurement Based on Impulse (Absolute Messverfahren oder Grund des Erkreuzens). A. Schmidt devel-

Ger. 1

oped a method which permitted calculation of the mechanical work produced by the detonation of an explosive. It is described briefly by A. Seutzbacher, Spreng- und Schießstoffe, Zürich (1948), p. 116.

Acetylene (Acetylen). See general section and the following references:
1) W. Kappeler, Advances in Acetylene Chemistry, PB Rept 1112 (about 1946).
2) W. Kappeler, Synthesis of laramonides for Polyamides or Acetylene Basis, PB Rept 25,553 (about 1946).

Active Sheet (Aktive Mantelgewebe) A type of sheet containing MG or PVC (nitrovinyl) together with inert ingredients was used by the Germans for some parachute explosives, such as Vetter-Waagig, etc. One of the earlier active sheets consisted of NG 15, rock salt 35 and Mg bicarbonate 50% but this was later changed to NG (with or without nitrovinyl) 12, rock salt 33 and Mg bicarbonate 55%. The composition of some other active sheets were:

Sheet	NG	NGC	NuCl	Mg bicarbonate	Kieselgahr
M ₁	10.0	-	35.0	55.0	-
M ₂	12.0	-	68.0	20.0	-
M ₃	11.0	1.0	87.0	-	0
M ₄	19.0	-	68.0	-	2.0

The sheeting operation was carried out automatically on the Sythe plant of W. S. A. A. on a modified Heppmann centrifugal machine, producing chippers weighing 70 gms to be sheeted with 55 gms of active sheeting material. Note: According to Seutzbacher (Ref 3) a sheet (Mantelgewebe) 25 mm in diameter and 3.5 mm thick, consisting of Mg bicarbonate 82% NG with NG 18 - 12%, indicates the composition of the gases of detonation from 2000' for a sheeted explosive to 400' C.

Note: According to T. Urbanski, Przemysł Chemiczny 6, 487, (1948), the active layer (sheet) was made in the form of a tube slightly larger than the cartridge of the engine charge. The cartridge was then inserted into the tube. When the cartridge was exploded, the combustible mixture layer (sheet) was dispersed and vaporized, thus forming a "cloud of salt" which prevented the ignition of fireproof oil and that which might be caused by the charge alone. (See also "Sheeted Explosives" in the general section).

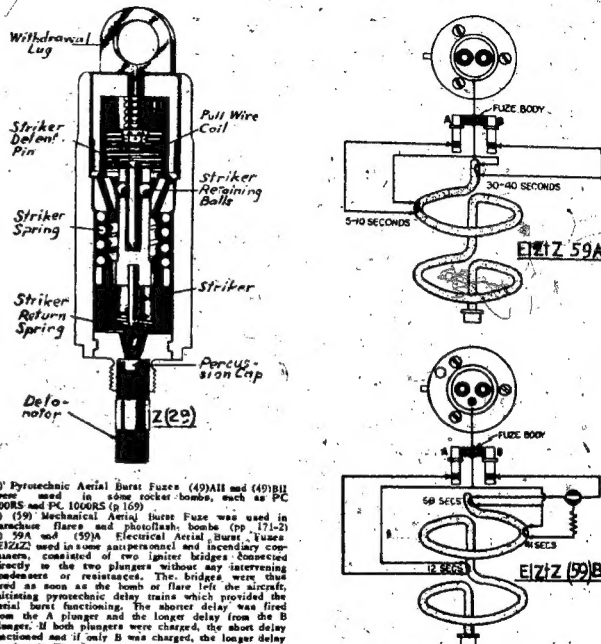
References:

- (1) O. W. Stichtel, PB Rept No. 925 (1945).
- (2) R. Ascher, PB Rept No. 65,877 (1946), pp. A-1/8 and A-1/11.
- (3) A. Seutzbacher, Spreng- und Schießstoffe, Zürich (1948), p. 92.

Aerial Burst Fuses are devices designed to function a bomb while still in flight. Following German fuses are briefly described in TM 9-1985-2 (1953), pp. 132, 166, 171-174:

- (1) (1) Mechanical Clockwork Fuse was used in SD 2A Bombsight bomb (pp. 132-3).
- (2) (2) Mechanical Aerial Burst Fuse, used in the LC 101 single unit parachute flare, consisted of a bakelite housing containing a clevis cap, withdrawal, top, safety spring, striker pellet guide, striker pellet, striker reset pin, firing spring, two ball detent, and a striker return spring. The withdrawal lug and the clevis cap were retained by a cord which was attached to the flare parachute. As the flare descended the safety spring was extended until it was tensioned sufficiently to withdraw the striker reset pin. The ball detent was then free to move forward and the striker pellet was forced by the firing spring to carry the striker into the percussion cap. At the end of an interval, the action pellet compressed the striker return spring. The flash from the cap ignited the delay element and, after the delay, the pyrotechnic mixture initiated the main charge of the bomb (pp. 166-7).

Ger 2



3) Pyrotechnic Aerial Burst Fuses (49) and (49) were used in SD 2A Bombsight bomb (pp. 132-3).

(1) (1) Mechanical Aerial Burst Fuse was used in parachute flares and photoflash bombs (pp. 171-2).

(2) (2) 59A and 59B Electrical Aerial Burst Fuses (EIZ 59) used in some parachute flares and incendiary containers, consisted of two igniter bridges connected directly to the two plungers without any intervening conductors or resistances. The shorter delay was fired as soon as the bomb or flare left the aircraft, initiating pyrotechnic delay trains which provided the aerial burst functioning. The shorter delay was fired from the A plunger and the longer delay from the B plunger. If both plungers were charged, the short delay functioned and if only B was charged, the longer delay functioned. The inner construction of both fuses was the same, but the 59A was twice as long as the 59B (p. 172).

(3) (3) Electric Aerial Burst Fuse (EIZ 59) used in SD 50 bomb and in some parachute flares, differed from the previous fuse by having three igniters instead of the conventional two, as to give a 1-second delay, was in such a position, as to give a 1-second delay. The other two igniters were under the B plunger and gave 41 and 58 second delays respectively. If the short delay was required, both plungers were charged. If a longer delay was necessary, only the B plunger was charged (pp. 172-3).

(4) (4) 69CL, 69L and 69E Electrical Aerial Burst Fuses (Pyrotechnic Delay), used in various bombs and containers, were cylindrical in shape and made of aluminum. The release from the plane, the igniter bridge light ignited the black powder. This in turn ignited the pyrotechnic mixture (in composition was given). On expiration of burning of the delay, the flash composition and the black powder pellet were

ignited, etc (pp. 174-5).

(5) (5) 79 and 79A Electrical Aerial Burst Fuses (Pyrotechnic Delay) used in parachute flares and photoflash bombs, resembled in appearance and action the 59 fuses (pp. 174-5).

(6) (6) 89, 89B, 89C and 89D Clockwork Aerial Burst Fuses are described on pp. 175-7.

The following aerial burst fuses are described in TM 9-1985-2 (1942), File Nos. 2314.9, 2324.91, 2324.92, 2344.91, 2342.9:

(7) (7) Mechanical Aerial Burst Fuse

(8) (8) Electrical Aerial Burst Fuse (See below)

(9) (9) Aerial Burst (Special) Fuse

(10) (10) Electrical Aerial Burst Short Time Fuse

(11) (11) Clockwork Aerial Burst Short Time Fuse. One of three fuses is described below:

(12) (12) (1) and (9) Electrical Aerial Burst (Short Time) Fuses, used in some parachute flares and in BG 30 photoflash bombs, were cylindrical in shape and contained a glow discharge tube, two condensers, a resistance, a

ELIZAB

ventilation. As it was difficult to cast-load Anatol 39 uniformly (without formation of cavities) in large caliber projectiles, G. Romer (Ref 5) used the so-called "bisquit" loading method. In this method, a projectile was filled alternately with pieces (pellets) of so-called "bisquit mixture A" (Am nitrate 50, technical Ca nitrate 25, PETN 10 and RDX 15%) and molten Anatol 39 at a temperature of about 80°. The resulting mixture formed no cavities on cooling. Its density at room temperature was 1.58, velocity of detonation 5600 m/sec. Translating this composition to 355cc for a 10 g sample one obtains a somewhat exact value (Stanchikov) (conservation of

Ammoncabini's See Werner-Ammoncabini's.

Ammoncabini's See Werner-Ammoncabini's.

Ammoniumnitrate (Ammoniumnitrate). A type of permissible explosive which may be considered intermediate between rhenolites and ammonium nitrate explosives. Table 1 gives the composition and properties of some of these explosives:

Table 1.

Composition (%) and some properties	Ammonium-nitrate (Ref. 1)	Ammonium-nitrate (Ref. 123)	Ammonium-nitrate (Ref. 124)
Ammonium nitrate	80.3	82	86.4
Potassium nitrate	5	10	-
Sodium nitrate	-	-	7
Nitroglycerin	4	3.5	2.0
Calcium carbonate	0.2	0.2	-
Charcoal	-	-	5.0
Carbohydrates (such as sugar, flour)	4.5	-	4.0
Coal dust	6	-	-
Alkali chloride	-	-	22.6
Wood meal	-	4	-
Oxygen balance	-	-	10%
Density	1.11	1.06	-
Velocity of detonation	3195 m/sec	3380 m/sec	-
Traust Test	55 cc	-	210 cc

References:

- 1) Marshall, 1957, p. 9, p. 493. 2) P. Naum, Nitroglycerin (1928), p. 434. 3) J. H. Ullmann, Encyclopaedia, v. 4 (1929), p. 780. 4) Davis, (1943), p. 352.

Ammoniumperchlorate (Ammoniumperchlorate). A type of powerful dynamic explosive containing a considerable amount of ammonium nitrate. Am nitrate 30.0, NG 67.0, collodion cotton 2.0, wood meal 5.0%; oxygen balance +1.55, density 1.44, Traust test value 485 cc, ball strength 21.0 mm, velocity of detonation 7000 m/sec, heat of explosion 1400 kcal/kg, temperature of explosion 2770°C.

This type of explosive was not very popular in Germany but was used in France and the USA [P. Naum, Nitroglycerin (1928), p. 493].

Ammoniumnitrate (Ammoniumnitrate). A type of permissible explosive, each part.

Ammoniumnitrate (Ammoniumnitrate). A type of permissible explosive. 1) DNT 7.8, Al 1.5-2.5, collodion cotton 0.2-0.7, diacetylenedinitrate (DNCN) 21-24, Am nitrate 61-65 and carbohydrates. See more than 1.5% (Ref. 1).

Notes: The Am nitrate may be replaced by Na nitrate to the extent of 8.5% of the entire explosive and the DNCN may be replaced by NG to the extent of 4% of the entire explosive. 2) Ammoniumperchlorate explosive prepared after WWI for use in Prussian mines. DNCN 54 which up to 3% of the total explosive was replaced by NG 28 to 32, collodion cotton 1 to 2, Am nitrate 45 to 50, alkali nitrate 10 to 15, a nitrocompound of collodion and/or naphthalene and/or diphenylamine 5 to 25. Specific heat 9 to 2% (Ref. 3).

3) Deutsche Ammoniumnitrate (DNCN), containing 15-20% of NG (which a mixture was called Nitrochlorite), 50, collodion cotton 3, nitrate of DNT and TNT 10, alkali nitrate 45, Am nitrate 10, wood meal 2, diphenylamine 45, velocity of detonation 6900 m/sec, Traust test value 400 cc, weight of plate at NTP 271 1/4 kg, heat of explosion 1101 kcal/kg, temp at explosion 2570°C, specific

pressure 8195 atm, brisance by the Kist formula 82,000 (Ref. 2 and 4).

Abbreviations: DNCN Diacetylenedinitrate; NTP Normal temperature (0°C) and pressure (760 mm).

[Compare with Ammoniumnitrate].

- 1) A. Marshall, Explosives, v. 3 (1932), p. 109.
- 2) P. Naum, Schiess- und Sprengstoffe (1927), p. 113.
- 3) P. Naum, Nitroglycerin (1928), p. 379.
- 4) A. Seebach, Spreng- und Schiessstoffe (1948), p. 86.

Ammonit (Ammonit) is described in the general section. The German method of manufacture of synthetic ammonit is described in BROS Final Rep. 1441 (1946).

AMMONIT (Ammonit) A type of ammonium nitrate explosive which has been known for many years and which exists in many varieties. Most ammonites were used as commercial explosives, but some of them have been used in military applications, chiefly as substitutes for hexanitroammonium for explosives based on organic nitrocompounds, such as TNT, or nitric esters (such as NG).

Many types of ammonites were known in Germany before WWI. For instance, Naum (Ref. 1) describes seven before, Berylls, and Detrop (Ref. 3) four types and Detrop (Ref. 3) on next page listing ammonites used during WWI for military purposes and see also under Commanded Explosives.

References:

- 1) P. Naum, Schiess- und Sprengstoffe, Steinkopf, Dresden (1927), pp. 119-121.
- 2) A. Seebach, Schiess- und Sprengstoffe, Barth, Leipzig, (1933), p. 246.
- 3) C. Berylls and K. Detrop, Schiessstoffe und Zündmittel, Springer, Berlin (1936), pp. 94-95.
- 4) O. F. Schindler, et al., General Summary of Explosives (1932), p. 104.
- 5) P. Naum, NG 925 (1945), Appendix 7, p. 77.
- 6) G. R. Rost, Report on Explosives, PBL Rep. No. 15, (1945), pp. 22-24.

Ammonium Nitrate Explosives

Ammonium Nitrate Explosives. See Ammoniumperchlorate.

Ammonium-Nitrat (Ammonium-Nitrat). A type of permissible explosive used after WWI, such as: a) Am nitrate 78.0, R nitrate 5.0, alkali chloride 8.0, meal 5.0, NG 4.0%; oxygen balance +11.85, Traust test value 200 cc, ball strength 21.0 mm, velocity of detonation 7000 m/sec, heat of explosion 1400 kcal/kg, temperature of explosion 2770°C.

b) Ammoniumnitrate (Ammoniumnitrate) 21-24, Am nitrate 61-65 and carbohydrates. See more than 1.5% (Ref. 1).

Notes: The Am nitrate may be replaced by Na nitrate to the extent of 8.5% of the entire explosive and the DNCN may be replaced by NG to the extent of 4% of the entire explosive. 2) Ammoniumperchlorate explosive prepared after WWI for use in Prussian mines. DNCN 54 which up to 3% of the total explosive was replaced by NG 28 to 32, collodion cotton 1 to 2, Am nitrate 45 to 50, alkali nitrate 10 to 15, a nitrocompound of collodion and/or naphthalene and/or diphenylamine 5 to 25. Specific heat 9 to 2% (Ref. 3).

3) Deutsche Ammoniumnitrate (DNCN), containing 15-20% of NG (which a mixture was called Nitrochlorite), 50, collodion cotton 3, nitrate of DNT and TNT 10, alkali nitrate 45, Am nitrate 10, wood meal 2, diphenylamine 45, velocity of detonation 6900 m/sec, Traust test value 400 cc, weight of plate at NTP 271 1/4 kg, heat of explosion 1101 kcal/kg, temp at explosion 2570°C, specific

pressure 8195 atm, brisance by the Kist formula 82,000 (Ref. 2 and 4).

Abbreviations: DNCN Diacetylenedinitrate; NTP Normal temperature (0°C) and pressure (760 mm).

[Compare with Ammoniumnitrate].

- 1) A. Marshall, Explosives, v. 3 (1932), p. 109.
- 2) P. Naum, Schiess- und Sprengstoffe (1927), p. 113.
- 3) P. Naum, Nitroglycerin (1928), p. 379.
- 4) A. Seebach, Spreng- und Schiessstoffe (1948), p. 86.

Ammonit (Ammonit) is described in the general section. The German method of manufacture of synthetic ammonit is described in BROS Final Rep. 1441 (1946).

AMMONIT (Ammonit) A type of ammonium nitrate explosive which has been known for many years and which exists in many varieties. Most ammonites were used as commercial explosives, but some of them have been used in military applications, chiefly as substitutes for hexanitroammonium for explosives based on organic nitrocompounds, such as TNT, or nitric esters (such as NG).

Many types of ammonites were known in Germany before WWI. For instance, Naum (Ref. 1) describes seven before, Berylls, and Detrop (Ref. 3) four types and Detrop (Ref. 3) on next page listing ammonites used during WWI for military purposes and see also under Commanded Explosives.

References:

- 1) P. Naum, Schiess- und Sprengstoffe, Steinkopf, Dresden (1927), pp. 119-121.
- 2) A. Seebach, Schiess- und Sprengstoffe, Barth, Leipzig, (1933), p. 246.
- 3) C. Berylls and K. Detrop, Schiessstoffe und Zündmittel, Springer, Berlin (1936), pp. 94-95.
- 4) O. F. Schindler, et al., General Summary of Explosives (1932), p. 104.
- 5) P. Naum, NG 925 (1945), Appendix 7, p. 77.
- 6) G. R. Rost, Report on Explosives, PBL Rep. No. 15, (1945), pp. 22-24.

Table 2.

Components and some properties	Designation of Ammonites											
	No. 7	43A	43B	44	43C	No. 7	41	45	No. 7	No. 7	No. 7	41-2
Am nitrate	42.0	46.0	56.0	45.0	46.0	50.0	50.0	55.0	52.0	50.0	-	-
Na nitrate	9.8	-	8.0	-	-	-	3.0	3.0	3.0	3.0	-	-
Ca nitrate, 4H ₂ O	-	8.0	6.0	10.0	8.0	15.0	15.0	10.0	7.0	15.0	-	-
Mg nitrate, 6H ₂ O	8.4	-	-	-	-	-	-	-	-	-	-	-
Guanidine nitrate	-	8.0	10.0	13.0	-	-	-	-	-	-	-	-
PETN	9.8	-	2.0	-	-	10.0	-	-	-	-	-	-
PH-Sala	-	-	5.0	-	46.0	-	10.0	10.0	10.0	-	-	-
RDX	30.0	8.0	7.0	-	-	25.0	20.0	20.0	25.0	25.0	-	-
Tetra-Sala	-	-	-	-	-	-	-	-	-	-	-	-
TNT	-	30.0	10.0	30.0	-	-	-	-	-	-	-	-
"Valuam" (emulsifier) (added)	-	0.3	0.3	0.5	-	-	-	-	-	-	-	-
Density (case)	-	1.58	1.61	-	-	-	1.53	1.50	-	-	-	-
Casting Temperature	-	104	105	-	-	-	108	112	-	-	-	-
Density of Fragments	41 m	-	38 m	-	-	39 m	-	40 m	41 m	-	-	-
Mining Effect	21 m ³	-	-	-	-	-	-	-	-	-	-	-
References	5	5	5	-	-	4	5	4	5	5	-	-

The composition given by R. Rost (Ref. 5, p. 22) (table 04).

* Ammonit 43C exploded in 1944 on a loading line and its manufacture was discontinued. It was reported that mixtures of TNT with guanidine nitrate were unstable.

Most of these mixtures were suitable for loading bombs, grenades and shells.

Am nitrate 90 and charcoal 10%. The mixture was compressed

in the form of perforated cylindrical pellets 4 to 5 cm long and 3 to 4 cm in diameter. The ignition temp of the compound was 160-165°, but at substances like iron rust, ZnO or CuO were present, the temp was lowered to 80-120°.

Notes: According to GOS 33-48, p. 7, the composition of Ammonipulver used during WWI was as follows: Am nitrate 50, NG (125N), 22, DEGN 22, hydrocellulose 5 and charcoal 1%.

References:

- 1) Marshall, 3 (1932), pp. 98-123. H. Herber, Chem-Ztg 59, 744-5 (1933).
- 2) Davis (1943), p. 49.

Ammonipulver (Ammonipulver) is described in the general section. Its manufacture in Germany at Stettin, Berlin and Witten plants is described in BROS Final Rep. No. 899 (1946).

Ammonipulver (Ammonipulver) is described in the general section. See Ammonium Nitrate Explosives, in the general section.

The German References on this subject include:

- 1) R. Exler, Ammonipulver, Steinkopf, Velt, Leipzig (1909).
- 2) P. Naum, Schiess- und Sprengstoffe, Steinkopf, Dresden (1927), pp. 114.
- 3) P. Naum, Nitroglycerin etc., Williams & Wilkins, Baltimore (1928), p. 423.
- 4) A. Seebach, Schiess- und Sprengstoffe, Barth, Leipzig (1933), p. 295.
- 5) K. Berylls, W. K. Detrop, Sprengstoffe und Zündmittel, Springer, Berlin (1936), pp. 99-100.
- 6) A. Seebach, Spreng- und Schiessstoffe, Baasche, Zürich (1948), pp. 86-88.

AMMONIUM (Ammonit) See under Bombs, Bullets, Cartridges, Fuzes, Grenades, Mines, Projectiles, Rockets and also in the following references:

- 1) Johnson, Jr. and C. T. Hawen, Ammonium, W. Morrow, N.Y. (1941).
- 2) Dept of the Army Tech Manuals, TM 9-45-2 and TM 9-45-3.

1965-5 (1965) Ministry of Technical Research, 1962

3) G.M. Tech. Ministry of Technical Research, 1962

4) W.H. Ewart, ibid, 1053 (1940) (20 mm Solchua CRA)

5) A.B. Schilling, ibid, 1168 (1942) (105 mm HE CRA)

6) A.B. Schilling, ibid, 1228 (1943) (88 mm APC HE CRA)

7) A.B. Schilling, ibid, 1238 (1943) (50 mm APHE SC CRA)

8) R.M. Dennis, ibid, 1242 (1943) (20 mm APHE CRA)

9) S.M. Dennis, ibid, 1245 (1943) (47 mm APC CRA)

10) A.B. Schilling, ibid, 1245 (1943) (47 mm APHE CRA)

11) A.B. Schilling, ibid, 1247 (1943) (75 mm APC HE CRA)

12) R.M. Dennis, ibid, 1248 (1943) (20 mm Inc. CRA)

13) A.B. Schilling, ibid, 1250 (1943) (50 mm APHE HE CRA)

14) R.M. Dennis, ibid, 1253 (1943) (37 mm APHE CRA)

15) A.B. Schilling, ibid, 1256 (1943) (20 mm HE SD CRA)

16) A.B. Schilling, ibid, 1258 (1943) (47 mm HE CRA)

17) A.B. Schilling, ibid, 1263 (1943) (80 mm Sn CRA for Inc)

18) A.B. Schilling, ibid, 1267 (1943) (50 mm APHE SC CRA)

19) A.B. Schilling, ibid, 1270 (1943) (50 mm HE CRA for Inc)

20) R.M. Dennis, ibid, 1271 (1943) (57 mm APHE HE CRA)

21) R.M. Dennis, ibid, 1272 (1943) (47 mm AP MB CRA)

22) R.M. Dennis, ibid, 1273 (1943) (50 mm APHE MB CRA)

23) R.M. Dennis, ibid, 1274 (1943) (50 mm APHE LC CRA)

24) A.B. Schilling, ibid, 1275 (1943) (20 mm AP Int. Loaded CRA)

25) R.M. Dennis, ibid, 1276 (1943) (75 mm HE CRA)

26) A.B. Schilling, ibid, 1300 (1943) (88 mm HE CRA)

27) R.M. Dennis, ibid, 1305 (1943) (50 mm HE SC CRA)

28) R.M. Dennis, ibid, 1314 (1943) (37 mm HE CRA)

29) R.M. Dennis, ibid, 1318 (1943) (57 mm HE LC CRA)

30) R.M. Dennis, ibid, 1320 (1943) (37 mm APHE MB CRA)

31) R.M. Dennis, ibid, 1326 (1943) (42/28 mm APHE CRA)

32) A.B. Schilling, ibid, 1329 (1944) (28/20 mm APHE CRA)

33) A.B. Schilling, ibid, 1331 (1944) (37 mm HE CRA)

34) A.B. Schilling, ibid, 1334 (1943) (75 mm HE CRA)

35) R.M. Dennis, ibid, 1340 (1944) (80 mm HE CRA for Inc)

36) R.M. Dennis, ibid, 1343 (1944) (75 mm HE CRA for Muz 40 gun)

- 30) A.B.Schilling, ibid, 1390 (1944) (28/20 mm HEHV CRA)
 31) A.B.Schilling, ibid, 1391 (1944) (88 mm HE LC CRA for Flak 41 gun)
 32) A.B.Schilling, ibid, 1392 (1944) (88 mm APC LC CRA for Flak 41 gun)
 33) A.B.Schilling, ibid, 1396 (1944) (37 mm HE HoC CRA)
 34) A.B.Schilling, ibid, 1421 (1944) (75 mm APC HE CRA)
 41) J.P.Wardlaw, ibid, 1422 (1944) (80 mm HE CRA for Mor) (Bouncing type shell)
 42) F.C.Hawerlak, ibid, 1430 (1944) (20 mm HE-T CRA for Mor)
 43) A.B.Schilling, ibid, 1434 (1944) (75 mm HE HoC CRA for How)
 44) A.B.Schilling, ibid, 1438 (1944) (75 mm HE CRA for How)
 45) A.B.Schilling, ibid, 1458 (1944) (50 mm HE LC CRA)
 46) F.G.Hawerlak, ibid, 1478 (1944) (20 mm HE HoC CRA)
 47) F.G.Hawerlak, ibid, 1481 (1944) (105 mm HE HoC 50 CRA)
 48) F.G.Hawerlak, ibid, 1487 (1944) (75 mm HE HoC CRA for recoilless gun)
 49) A.B.Schilling, ibid, 1488 (1945) (150 mm HE HoC CRA)
 50) J.P.Wardlaw, ibid, 1490 (1945) (75 mm HE HoC CRA for Pak 40 gun)
 51) F.G.Hawerlak, ibid, 1496 (1945) (105 mm HE HoC Type C-L shell CRA)
 52) F.G.Hawerlak, ibid, 1503 (1945) (75 mm HE HoC CRA for KwK 40 gun)
 53) F.G.Hawerlak, ibid, 1508 (1945) (100 mm APC HE CRA)
 54) F.G.Hawerlak, ibid, 1516 (1945) (88 mm APC HE CRA for KwK 41 and Pak gun)
 55) A.B.Schilling, ibid, 1522 (1945) (150 mm HE CRA, separate loading)
 56) A.B.Schilling, ibid, 1529 (1945) (150 mm HE A/C CRA w/BD fuse)
 57) F.C.Hawerlak, ibid, 1540 (1945) (75 mm HE HoC CRA for short barrel mark gun, KwK 38)
 58) F.G.Hawerlak, ibid, 1551 (1945) (150 mm How CRA)
 59) F.G.Hawerlak, ibid, 1552 (1945) (210 mm HE CRA)
 60) A.B.Schilling, ibid, 1559 (1945) (88 mm HE, serrated shell for Flak 18 gun)
 61) F.C.Hawerlak, ibid, 1575 (1945) (150 mm CP shell and cartridge case with poppet of Russian origin)
 62) A.B.Schilling, ibid, 1577 (1945) (240 mm HE shell with PD and BD fuses, charge case and poppet)
 63) A.B.Schilling, ibid, 1578 (1945) (75/75 mm HE CRA for tapered bore Pak 41 gun)
 64) A.B.Schilling, ibid, 1579 (1945) (75/55 mm APC CRA for tapered bore Pak 41 gun)
 65) A.B.Schilling, ibid, 1582 (1945) (100 mm HE CRA for Mor)
 66) A.B.Schilling, ibid, 1604 and 1605 (1946) (105 mm rocket assisted HE shell)
 67) A.B.Schilling, ibid, 1606 (1946) (128 mm rocket assisted HE shell)
 68) A.B.Schilling, ibid, 1607, 1608 and 1609 (1946) (130 mm rocket assisted HE shell)
 69) A.B.Schilling, ibid, 1610 (1946) (150 mm rocket assisted AP shell)
 70) A.B.Schilling, ibid, 1903 (1954) (30 mm HE and HoC shell for the A/C Mt-108 cannon (Confidential))
 71) Anon, Enemy bombs and Fuses, War Dept TM-E9-1983 (1952)
 72) Anon, Enemy War Materials Inventory List, Ammunition, Supreme Headquarters AEF (1945)
 73) Anon, Recognition Manual of German Ammunition, Supreme Headquarters AEF (1945)
 Note: All Pictorial reports except No 1903 are unclassified
 Abbreviations: AA Antiaircraft; AC Aircraft; A/C Aircraft; APC Armor-piercing; A/P Anti-personnel; BD Bouncing

detonating; C Capped; Chem Chemical; CP Concrete-piercing; CRA Complete round of ammunition; Flak German designation of Antiaircraft; HC High capacity; HE High explosive; HoC Hollow (shaped) charge; How Howitzer; HV Hyper-velocity; Inc Incendiary; KWR German designation of Tank-Gun; LC Long case; LQ Long ogive; M8 Mounchick; Mor Mortar; Pak German designation for Antitank; PD Point-detonating; SC Short case; SD Self-detonating; Sm Smoke; SO Short ogive; T Tracer.

"Amores" (Toy Pistol Cap). Due to the shortage of fulminate for some hand grenades, Amores manufactured by Ferdinand Wicke, Tupper-Leno and by Blumberg & Co, Linum Ltd, Düsseldorf combined; K chlorate 0.75 g, RbO₂ phosphorus 12.5 g, SnO₂ sulfur 8.9 g and chalk 13.3 g, 3.7%
 Reference: RUS Final Rep 1311 (1947), p. 2-4.

Amorger. One of the early aluminum explosives: Al 5.5, Am nitrate 84.5, K nitrate 1.5, charcoal 8.0, Ba nitrate 0.5% (L. Midard, Min Artil Fr 72, 966 (1948)).

Antioxiolite (Antioxiol Cap). Due to the shortage of brass during WWI, the Germans used zinc and zinc-coated iron caps. They were filled with TNT as the base charge and compressed silver fulminate as the primary charge. The assembly was called Antioxiolite. [P. Warden, Schies- und Sprengstoffe, Stuttgart, Dreyden (1927), p. 185].

Antitank Wind Gun. See Wind Gun.

Antitank Fuse (Antitank Fuse). Such as AZ (204) was a mechanical impact fuse with a safety timing period of 10 seconds provided by the clockwork gear train. There were two striker systems incorporated: an impact striker system to operate on impact and an antitank-striker system to function in case there was any distortion of bomb or fuse-pocket on impact. The two striker systems were located at opposite ends of the fuse as opposed to a flash channel about 200 mm long. This fuse, as well as the AZ (24), are described in TM 9-1985-2 (1953), pp. 13-9. They were used in bomb SC 2500 kg. (See illustration on next page).

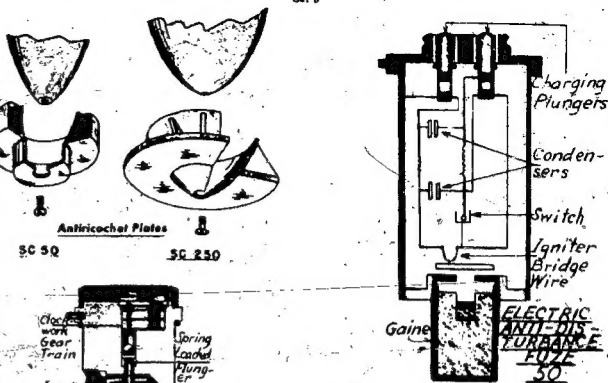
Antishock Fuse (Electrical) was a device designed to function if disturbed after the bomb dropped from the air had come to rest. One type, the 30, consisted of a cylindrical case containing an electrical circuit (two condensers, two resistors, a representative ball-bearing switch and bridge wire of primer) and two charging plungers. The base of the cylindrical case was threaded to receive a gasket. Before dropping the bomb, the electrical circuit from the plunger was connected through the charging plungers into the charging condenser. During the flight the charge slowly leaked through a high resistor in the lifting condenser. If after the bomb had come to rest it was subsequently disturbed, the triaxial circuit caused the lifting condenser to be ignited, the primer, initiated the booster and detonated the HE charge of the bomb. This also took place when one or both charging plungers were removed, thus shunting the current from the condenser by-passed the switch. [TM 9-1985-2 (1942), file No 2327]
 Another antishock fuse, the 32, was Y was much more complicated. Its description is given in TM 9-1985-2 (1953), pp. 183-6.
 (See illustration on next page).

Antiflight Igniter. See Item 1 and L under Igniter.

Antipersonnel Devices. See Pyrotechnic Antipersonnel Devices.

Antitank Bomber. See Senger-Berck Missile.

Anti-Rocket Plume. Circular shaped metallic devices attached to the nose of some aircraft bombs intended to prevent ricochet when striking at an angle of obliquity against very resistant targets (such as armor), or to prevent undesirable penetration into more resistant targets (such as concrete or wood) when striking them at an angle close to normal. (See illustration on next page and also under Kofing) [TM 9-1985-2 (1953), p. 4].



Antishock Fuse (Mechanical) were designed as protective devices to prevent withdrawal of regular time fuses from bombs. Three types of such fuses are described in TM 9-1985-2 (1953), pp. 177 & 179-181: Xuz 40, Types I, II and III. The type I fuse consisted of a cylindrical body with a central opening in the upper surface to receive the gain of the time fuse which it protected, and a second gain which was threaded into the base of the Zuz 40. An attempt to withdraw the fuse would cause the steel ball (below the detent spring) to be displaced, thus allowing the striker to hit the detonator. The resulting flame was transmitted through a small channel to the booster, and the bomb was exploded. To prevent the withdrawal of the Zuz 40 when the time fuse was removed, spring-loaded knife edges were placed in the upper part of the device. (See illustration on next page).

"Annie Antia" or "Loppled 280 mm Railroad Gun, Model 5 (See under Weapons).

Armored Cars are described in the following references: 1) G.B. Jurek, "Achungs Panzer", Great Oaks, RD 1, Aberdeen, Md (1948)
 2) D.F. von Senger u. Esterlin, "Taschenbuch der Panzer", Lehmann, München (1954) (See also under Panzer).

Art (Azur). A mining explosive reported to be made by VEB Sprengstoffwerke, Gschwitz, its approximate composition is NG, pyroxilin, sawdust, TNT and inorganic salts.

Armhook (Needle Point) Projectile, such as 5 cm Pakgr 40 for 5 cm Pak, was a 30 mm AP Proj which consisted of the following components: a pointed tungsten carbide core cemented to a steel body which had forward and rearward flanges; a plastic armhook shaped head covered with a sheet steel ballistic cap and a cruet assembly. The forward flange acted as the rotating band, while the rear flange acted as the bourrelet.

The Pak 40 was fired from a normal gun. On striking the armor, the ballistic cap, the head and the body with

Ger 9

ward assembly were shattered thus leaving the tungsten carbide core to penetrate the armor.

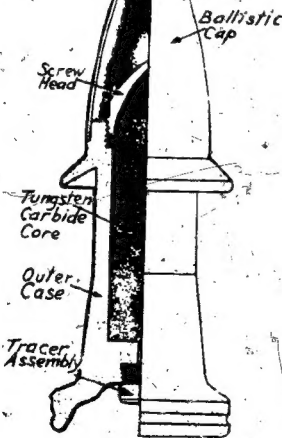
By employing the strawhead design, the weight of projectile was about half the conventional PzP (HEAP) shell. Due to this lightness, it was possible to develop very high velocities and high armor penetration at short ranges. The projectile was, however, very inaccurate at long ranges and the penetration hole was small in comparison with the gun caliber. (See also Tapered-Bore Gun Projectile).

There were also 37 mm (3.7 cm PzP/PzF 40), 47 mm (4.7 cm PzP/PzF 40) and 75 mm (7.5 cm PzP/PzF 41) strawhead type AP projectiles.

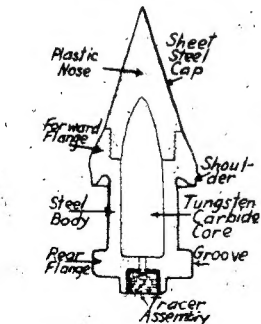
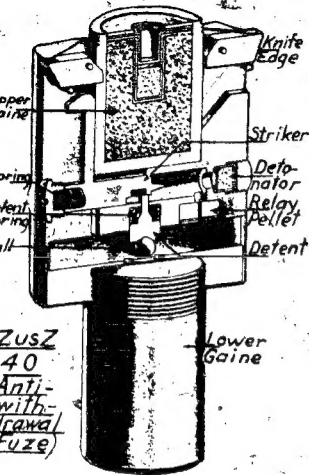
References:

- 1) E. Engelberg, Ordnance Sergeant May 1944, pp. 311-312
- 2) Same, Technical Manual TM 9-1985-3, pp. 373, 376-2.

**75 MM
AP PROJECTILE
WITH TUNGSTEN CARBIDE CORE
(ARROWHEAD)**



Arrow Projectile ('Pfeilenhaube Geschoss') is a slender, very long, fin-stabilized subcaliber projectile fired from a smooth-bore gun at supersonic velocity. Its development which is described in Ref 1 may be considered as one of the outstanding German achievements of WW II. Some of these projectiles were used in the attack on the Maginot Line and were successful in penetrating concrete. Their subsequent use was confined to the Russian front.



ARROWHEAD PROJECTILE

Ger 10

Artillery Ammunition (Complete Round). See under Grenade.

A5. Abbreviation for Ammoniumsalpetersaure (Explosives based on ammonium nitrate) [Weichert (1953), pp. 35, 375.]

A5-3. One of the German priming (lighting) compositions used during WW II in some electric fuzes. It contained red lead 77, silicon 19, NC suspended in acetone 4 [PB Rep. No. 95 613 (1947), Section 1].

ASN. See under Unterschweinskeigroffe.

A-Stoff (Liquid Oxygen) is described in the general section. It was used in some liquid propellants for guided missiles such as the A-4 (V-2), Teufel and Wasserfall.

Reference: Gollay, Rockets and Missiles, CIOB No. 28-36 (1946), p. 3.

Note: According to CIOB 33-13, p. 20, the A-53, which means Artillery School composition 3 (Artillereschule 3), was an incendiary composition prepared by mixing 75 parts of red lead with 25 p of silicon made into a paste with NC jelly.

Assisted Take-Off (ATO) Units. See under Rocket.

Australit (Australite). A type of mining explosive similar in composition to Ammonal and Dynamite.

Typical compositions are given in the following Table 3a.

Table 3a

Composition (%) and some properties	Australit 1	Australit 2	Australit 3	Australit 4	Australit O N
Ammonium nitrate	84.5	80.0	79.0	68.3	80.0
TNT + DNT	7.0	12.0	-	-	-
Vegetable meal	1.0	3.0	-	-	-
TNT + DNT + meal	-	-	17.5	27.7	20.0
Charcoal	1.0	1.0	-	-	-
Paraffin oil	2.5	-	-	-	-
Nitroglycerin	4.0	4.0	4.0	4.0	-
Oxygen Balance, %	-	-	+2.5	-	+6.3
Transit Test, cc	-	-	390	-	375
Pb Block Crushing, mm	-	-	16.2	-	16.0
Sensitiveness to Initiation (requires)	-	-	No 1 Cap	-	No 3 Cap
Propagating in 30 cm Cartridge	-	-	12.0 cm	-	80 cm
Velocity of Detonation m/sec	-	-	5400	-	4900
Density of Cartridge (test of Explosion, kcal/kg	-	-	1.09	-	1.03
Temperature of Explosion, °C	-	-	2170	-	2270

*See Propagation of Explosion in Cartridges, described in the general section.

References:

- 1) A. Marshall, Explosives, I, (1917), p. 397
- 2) P. Neomin, Nitroglycerin (1928), pp. 423 & 426.

Athyphenylurethane (Ethylphenylurethane) was used as an ingredient of some smokeless propellants (as a stabilizer-gelatinized) [PB Rep. No. 11,544 (1943)].

Auroral Impulse. See T-Stoff.

Ausbauchungsprobe (Expansion Test). See Transl Lead Block Test in the general section.

Ausschwitzungsprobe (Sweating Test). See Exudation Test in this and in the general sections.

Axeton (Acetone). See general section.

Azide (Azides) are described in the general section. (See also this section under Bleisatz).

Amaloid (Stichtstoffverwitterung) (Hydrazine Acid), See general section.

A. A. vehicle 12' long, 6' wide and 4' high provided with 6 cylinders engine (in the rear), 2 radio sets, 1 type for the driver. After loading the vehicle with some demolition charges, the driver test the car (max speed 30 mph) as follows: is possible on the target material for destruction (such as a buried mine, land block, pillbox, bridge, etc.) stepped the demolition charges, but the time fuse and then marked back. These vehicles were easy targets for the Allies' artillery.

Reconnaissance, Army, Field Artillery, J. 34, 505 (1944).

See (Bachem 247 Model), See Helmer No 3494 and 3495.

Explosive: One of the mining explosives: Am nitrate 85 and TNT 15%. (L. Colvin, High Explosives (1918), p. 249.)

"Butor", See "Horus" named vehicle listed under France.

Ballistische Beständigkeit (Ballistic Stability): See general section.

Sulfate as WFC/99 (Wolfschmelze/99) (Cube Powder of 1999) (Ballistics): Dark gray precipitate consisting of equal parts of Mg and cellulose coarse together with 0.5% to 1% of DAP and vasoline. It was adapted in 1889 by the German Navy. Last amateur competition was conducted in 1897 and 1900, under the designation of RPK-97 and RPK-00, when RPK was known for Rhamphor (tube powder) [Marschall, v. 1 (1917), p. 565].

Bandholz Pulver, was prepd by compressing the Schmelze Pulver, last sample of high density [L. Colvin, Titled: The Heilbronn Explosives, Mainz, (1907), p. 469].

Hampe's Topf (In Holz gefüllte Reithöhle). See general section.

Ble (Bee). One of the experimental units (See under France).

Bombol See general section.

Borax (Borax). Under this title, Daniel, Dictionnaire (1902), p. 57 gives a mixture of 8 parts of black powder with 2 parts of borax. It was used in the 1860's in large caliber guns.

Bromwasser (Cement). See under Cellulose in the general section.

Bromwasser (Overpowered or Malicious Mind): Several lead mines used by the Germans during WW II were made from iron not specially designed for mines. For instance, Schmelze No. 1, A/P, was improved from composed 50 mm water shell. Several important lead mines had described on pp 279-83 of M 1-1985-2 (1953).

Bullbomg, See Bommer Charge and under Ignition.

Bullit (Bullit). One of the Sprengel type explosive, it was also used in England and other countries (See in the general section).

Burr Barrel. See Krummholz.

Beschussungsgegenstände (Observation Round): Fired round with a projectile which had a core of R.E., a fuse in the central portion and a phosphorus filler in the base.

The purpose of this round was to indicate the target location of a hit by means of a puff of smoke (produced on ignition of the phosphorus).

Reference: A. Dore, Ordnance Sergeant, Dec 1943, pp 357-61.

Beryllite B. According to L. Moland, Météor. Air Fr 22, 1965 (1948), the Beryllite 'B' is one of the older aluminum explosives: Am nitrate 35%, DNT 5, Mg 5, cellulose cotton 4.5, Al 5 and cellulose 5%. Later the French lead block explosion, was (modified Treussart) in 125, which the value for pyritic acid as 100.

Bergmischung (Bergmischung): A smoke-producing mixture composed of 2 parts of zinc dust and 3 parts of berolichowite [U.S. War Dept Tech Manual, TM 30-506 (1944), p. 23].

Bergmischung-Junk Stability Test. See general section under Stability Tests.

Bewehrter oder Verformender (Tamping or Seaming). See general section.

Beschussungs (Shooting Test, called in the U.S.A. Rifle Ball Test): It is similar to the U.S. test described in the General section. The German test is conducted according to: Southworth, Spang and Schickeloth (1948), p. 121 by firing a standard rifle bullet (in a distance of 25 meters).

Bleisch Continuous Process for the Production of Nitroglycerin and Nitrocellulose as used in the Dynamit A-G, Schmelze Pulver is described by Dr. V. B. Linder, D.B. Chapp, ROR Final Rept 1842 (1946) (See also under general section).

Bleisch Explosives: Several compositions were patented by C.E. Biche) at the end of the last century, among them: a) NG 100 pulverized with 10 p of subchlorinated, b) N nitrate 90-100 p mixed with 10 p of nitrocellulose and 10 p of subchlorinated (u), c) Am nitrate 85 p mixed with 8 p of TNT and 6 p of lead or starch.

Reference: Daniel, Dictionnaire (1902), p. 67-8.

Big Bertha Gun. See general section.

Bismuth (Bismuth). A type of permittible explosive containing large amounts of sodium bicarbonate and small amounts of NG, patented by W.A.S. & Co before WW II. These explosives, although they contained a large amount of NaHCO₃ and a small amount of NG, were very easy to ignite. However, considering that 80% NaHCO₃ and as little as 20% NG could still be ignited by ordinary blasting caps.

The following are the composition and properties of a) of the bismuthine NG 15, NaHCO₃ 50 and NaCl 35%; temp of ignition 400°, value of 1000 ft of explosion 162 kcal/kg; 4:1:35, Tminal test value 50 cm for a 10 g sample, specific pressure 610 atm 1/kg, nitrogen value (Kapt) B = 4 (in (pp) units) (value of det) $\times 10^{-4}$ = 2.06, gas test value (Dynamisch) beschussungsgegenstände (Schlagempfindlichkeit) 60 cm, required for initiation at least a No 2 blasting cap, volume of gases evolved on explosion of 1 kg in 258 l at 20 and 760 mm Hg (O₂ in vapor phase). Composition of gases: CO₂ 66.1, H₂O 43.2, N₂ 5.2, and C 1.2%.

Note: When a more brilliant explosion is desired, the amount of NG is increased, the amounts of NaCl and NaHCO₃ are decreased and some fuel and oxidizer are incorporated.

The following mixture may serve as an example of such an explosive: NG (slightly calcinated) 30, NaHCO₃ 40, NaCl 12, wood meal 4.5 and NaNO₃ 13.5%; temp of explosion 1400°; vel of deton 4000 m/sec. d 1/4, Trauzeit value 124 cm for a 10 g sample, gas test value 30 cm; could be initiated by a No 2 blasting cap.

The bismuthine were comparatively expensive, but they proved to be very safe for use in gaspans or dusty coal mines.

Reference:
C. Beyling, A. K. Dreppel, Sprengstoffe und Zündmittel, J. Springer, Berlin (1936). Reproduced by Edwards Bros, Ann Arbor, Mich., pp 145-146.

Bismut Mixture A. See under Amaloid 39.

Black Powder. See Schwarzschnitzpulver.

Bleaching Cap. See Detonators.

Bleaching Gelatin. See Sprenggelatine.

Bismuthpulver oder "B" Pulver (Lead Powder or Flake Propellant): According to Stenbacher, Spreng- und Schmelze (1948), p. 94, it was prep by colliding a mixture of 3 parts of gun cotton (Schmelze) of N content minimum 15% and 1 p of soluble NG (Nitroindium) of N content 15%. After incineration into mixture 0.5% of the stabilizer (DPA) and 1% of flash-reducer (Na oxalate), the mass was flaked and dried; the resulting lumps (which were 0.5 mm thick and had a surface of 1.5 mm²) were surface-coated with crystalline and finely pulverized graphite in order to make them progressive burning.

Blende (Lead Aside) (L.A.). See general section, under Asides. It was used in Germany in some priming and initiating compositions. L.A. was prep in Germany during WW II from sodium aside and lead nitrate in the presence of dextrin, in the following manner:

- Fifty liters of water containing 1.5 kg of sodium aside was added slowly to 60 l of an aqueous solution containing 5 kg of PbO and 0.15 kg of dextrin, preheated to 60° and stirred by air. After adding the first 5 liters, there was a pause of 5 minutes. The remaining 45 l was added during the next 45 minutes, and the stirring was continued for 15 minutes, while the mixture was cooled by means of cold water circulating through the jacket.
- Following this, the reactor was tipped onto a filter and the L.A. retained on a filter cloth made of horse hair. Suction was applied.
- After rinsing the L.A. with several portions of water, it was placed on sheets of paper attached to frames and dried to a moisture content below 0.1%. Drying was done by blowing air (at 48 hr to 4-500° through the chamber containing the frames).
- After cooling to 200° the contents of each sheet were transferred to a spherical cup-shaped dish. The desired amount of dried L.A. was added to the same dish, which was then sent to detonator manufacturing plant. (Yield was about 3.3 kg per batch).

In order to destroy any L.A. remaining in the mother liquor, about 5 liters of nitric acid (50%) and about 1/2 l of concd H₂ nitric acid were added per batch of L.A.

Reference: Pt Rept 55,613 (1947), Sections O & P.

Note: According to L.M. Sheldon, "Manufacture of Initiating Explosives", GIDP 1938, p. 718, the manufacture of L.A. at the Woburn-based Plant of Dynamit A-G was conducted in a large, well-ventilated, stainless steel, round-bottomed cylindrical reactor. Jacketed for circulation of heating or cooling water or brine. Agitation was conducted by one motor located along the top of the reactor, as shown on the attached drawing. This agitator could be raised or lowered as required to provide the most efficient stirring. The agitator could be raised or lowered as required to provide the most efficient stirring. For discharging the contents of the reaction vessel the agitator shaft was fitted with the handle which was then tilted by a control lever. The agitator shaft was supported by two bearings. A stock solution of 10% by weight of lead nitrate and 1% to 5.0% by wt of sodium aside were kept in large vessels placed higher than the reactor in order to secure the flow of liquids by gravity. The correct volume for each precipitation charge was obtained by the use of calibrated glass bottles.

Flow rates were controlled by manually operated stopcocks. Before pouring with precipitation the alkalinity of the sodium aside solution was checked by titrating with normal sulfuric acid. It was acceptable for use, 50 ml of aside solution required 8 to 10 ml of acid to reach the phenolphthalein end point. If the aside was not sufficiently alkaline, some silver sodium hydroxide was added in the stock aside and the titration was repeated. Oxidation was checked by making the stock solutions.

In carrying out an individual precipitation, the volume of sodium required to give 4.3 kg of actual lead nitrate (500 l when using a 95% aside) was drawn from the supply tank and measured in a graduated glass bottle from which it was transferred to the reactor. (This amount of first material was about 0.8% in excess of that theoretically required.) After heating the aside to about 50° some dilute sodium hydroxide was added until the aside became neutral to methyl orange, as determined by a spot plate test. After neutralization, 150 g of potato dextrin (which had previously been dispersed in warm water) was added to the aside. The correct volume of sodium aside solution to give 15 kg of actual material (500 l when using a 95% aside) was measured in a calibrated glass bottle (the which was discharged through an adjustable stopcock into the reactor, and constantly stirring the solids and maintaining it at 50°. The rate of flow was controlled by a stopcock.

Na aside solution was added at a fairly uniform rate over a period of about 1 hour. After addition of the Na aside solution had been completed, the agitator was stopped, the lead aside allowed to settle and the mother liquor decanted by tilting the vessel. After giving one dilution wash first in the reaction vessel, it was tilted and the precipitate transferred by means of a pig or into a large container. The mother liquor was a natural drainage filter. After finishing the lead aside with three displacement type washes, the sludge was folded over the sides and the mother liquor was decanted. The mother liquor was carried to the average line. The yield was about 3.3 kg of detonated lead aside.

A sample of each batch was sent to toxicology where the crystals were examined microscopically and compared with authentic standards. Two parts of aside was dried and its loading density was determined.

For destruction of unwanted L.A. it was treated successively with a 15% H₂SO₄ acid solution and an 8% Na aside solution (1948).

Note: Crystalline structure of L.A. is described by G. Pfefferkorn in the Zeitschrift für Naturforschung, 3, 164 (1948).

According to W. Schneider, Sprengtechnik No 10-11, pp 185-196 (1952) and Explosivstoffe No 1-2, pp 1-10 (1953), technical L.A. (purity 92-94%) used in German Sprengkapsel (A and Sprengkapsel B) becomes dead-pressed if the loading pressure exceeds about 1800 kg/cm² (about 12,800 psi) depending on conditions. Perfectly dry L.A. can stand higher pressures without being dead-pressed, but L.A. can



The image contains several technical drawings of bombs and incendiary devices, with labels for various components:

- Top Row (Left to Right):**
 - 500-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 500-lb Bomb.
 - 1000-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 1000-lb Bomb.
 - 5-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 5-lb Bomb.
- Middle Row (Left to Right):**
 - 500-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 500-lb Bomb.
 - 1000-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 1000-lb Bomb.
 - 5-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 5-lb Bomb.
- Bottom Row (Left to Right):**
 - 500-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 500-lb Bomb.
 - 1000-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 1000-lb Bomb.
 - 5-lb Bomb:** Labels include Tail Fin, Bomb, Main Fuse, Main Fuse, Main Fuse, Explosive Charge, Subincendence Lin, Incendence Lin, Fuse Pin, Explosive Charge, and 5-lb Bomb.

2) Kist-Metz, *Chemische Untersuchung*, Vieweg, Braunschweig (1944), p. 165.

2) Kist-Metz, *Chemische Untersuchung*, Vieweg, Braunschweig (1944), p. 165.

(See also Colored Smokes in the general section).

eration was followed by addition of a weighed amount of NG-diisoglycol mixture, while continuing the hand stirring. The resulting gelatin was allowed to stand for 1 hour.

ter: For Am nitrate-type explosives, the plasticity was sometimes controlled by adding a solution of "gelose".

4) The pan was removed to another building where it was placed under the outlet funnel of a sieve through which the usual solid components of dynamite (such as Al or Na nitrate, TNT, wood meal, etc.) were fed. These components were previously pulverized and dry blended in another building. While the addition of the solid ingredients took place, the mass in the pan was stirred by means of a planetary stirring mechanism which could be lowered or raised as desired. Kneading time was usually about 20 minutes.

49: Several types of mixers (blenders) were used, such as the Deisawerke, Vetrig, McRobert and a modified Pflaider.

c) The thoroughly kneaded mass of gelatin and of solid components was removed by a wooden hand spade into wooden transport boxes to be carried to the canuliding plant.

test: German permissible explosives, were usually white in color, while the non-permissible were colored red by addition of caput mortuum (FeO) in the mixing stage. d) Cartridgeing was done either by fully automatic machines (such as the system of Niepmann & Co., Gera, Germany) or by semi-automatic machines (such as the system of Breuninger). The Breuninger machine (made entirely of brass) consisted of a cooidal casing through which passed a horizontal feeding screw. The grainized mass was introduced into the machine by hand through

the filling funnel. A paper cartridge was placed at the narrow end of the conical casing. After a cartridge was filled, it was removed by hand and the open end crimped. The diameter of a cartridge was 22, 23 or 30 mm. After packing these cartridges into a box (36, 35 and 26 cartridges per box, respectively), this box was wrapped in paper and dipped in paraffin. For shipping, 10 boxes were packed in a case.

⊗ Permissible explosives were mechanically shattered with a "reactive shaker" consisting of NG 12, NaCl 33 and NaHCO₃ 55% (NaHCO₃ 55% is usually 50% NaHCO₃ and 5% Na₂CO₃). The shaker was NG 15, NaCl 35, NaHCO₃ 50%. The shaker contained 33 s and the output itself 10 s.

Powder type explosives: To this type belong explosives which contained small amounts of NG (such as 4%), as diiodine gases and were pulverulent. The mixing of the components was done in a suitable type Werner-Pfleiderer mixer which consisted of a brass trough provided with 4 horizontal brass stirring rollers running in opposite directions.

a) The weighed amounts of the solid components (such as

Am, Co or Na nitrate, TNT, wood meal, etc. were mixed in a Werner-Pfleiderer blender, the liquid DNT, NG, NGC etc. were added and the mixture kneaded for 15 minutes.

b) The kneader was then tilted and the mixture discharged into wooden casks to be taken to the loading place.

c) In the case of explosives such as Calcium, the mass could be immediately cartridged, but with Donasit the mass had to be left overnight in storage before cartridging.

Table 8 gives some typical German Commercial Explosives manufactured before and during WW I.

Table 4

Ingradients and some properties	Ammonium I (1932)	Donastr I	Dynastr I	Gelatin Donastr I (1936)	Wetter Donastr A (1936)	Wetter-Nobelstr B (1932)	Wetter-Nobelstr B (1932)	Wetter-Wagstr A B	Wetter-Wagstr A (Permissible)
NC(Nitrocellulose)	4.0	4.0	63.0		6.0	21.4	29.2	30.0	20.8
NGc (Nitroglycerol)				22.0					4.0
NC(Nitrocellulose)			2.0	0.8		0.6	0.8	1.0	0.7
TNT(Tri-nitro-toluene)	6.0	32.0		5.0	2.0				0.5
DNT(Di-nitro-toluene)(liquid)	6.0	2.0		6.0		2.0			0.5
An nitrate	80.2	79.8		55.0	72.0			29.5	30.5
No nitrate			26.7	10.0		32.0	26.5		80.5
Wood, meal	3.5	2.0	8.0	1.0	2.0	1.0	0.5		0.3
Rock salt(NaCl)					18.0	56.5	40.0	59.0	59.5
Caput mortem (Fe ₂ O ₃)	0.3	0.2	0.2	0.2					13.0
Gelose (Carra-gine moss)								0.5	0.7
Talc									0.3
50% Ca nitrate solution						2.5	3.0		
Transit Test, cc	370.0		385.0	380.0	220.0	209.0	185.0		
Lead Bloch	17.5		23.0	20.0	10.5	6.5	11.0		
Compression, mm									
Veloc of Deton., m/sec.	4800 (as 4.1-10)		6350	6150	3000 (as 4.1-10)	5750	5650		
Cartridge Density, g/cc	1.67		1.55	1.53	1.06	1.66	1.69		
Gey Test, cm	6.0		10.0	10.0	8.0	6.0	7.00		
Charge Limit, g					600		700		
Oxygen Balance, %	+0.06		+5.0	+3.68	+10.4	+4.98	+6.15		
Heat of Explos., kcal/g	996.0		1291	1029	516	642.0	508.0		
Gas Volume, l/kg	904.0		609.0	806.0	772.6	536.0	500.0		

References:

1. O.W. Stickland, <i>General Summary of Explosives Plants</i> , PB Rept 925 (1945), p 69	Explosives, B I O S Final Rept 833, Item 2
2. R. A. Schrock et al, <i>Investigation of German Commercial Explosives</i> , PB Rept 6, 877 (1946), pp A 1/8 and A 1/11.	3. R. A. Schrock et al, <i>Investigation of German Commercial Explosives</i> , PB Rept 6, 877 (1946), pp A 1/8 and A 1/11.

Complete Round of Artillery Ammunition. See under *Grenat*

Composition A (Comp A) A mixture of RDX 90-97 and Montan was 10-35, similar in properties to Comp A used in the U.S.A. during WW II and described in the general section. German uses of Comp-A were in boosters, sub-boosters and as a filler in some grenades and shaped charges. (See also-Filix No 86, No 91 and No 92).

Reference: Allied and Enemy Explosives, Aberdeen Proving Ground, Md. (1946), p 122.

Composition B (Comp B) (Cyclotol) A mixture of RDX and TNT in various proportions similar to Comp B described in the general section. Some of the compositions contained small amounts of wax. Comp B was used by Germans, during WW II for filling shaped charge shells, grenades, rockets and some demolition charges. Pellets of Comp B embedded in TNT were used in 4000 kg bombs.
(See also Filler No 18 and Filler No 95).
Reference: Allied and Enemy Explosives (1946), p 124.

Composition: C. A plastic explosive similar in properties

Detonants A (Detonanten A). One of the Favier type explosives: Amtrazene 90.8, K bicarbonate 2.2, naphthalene 6.5, carotene 0.05, vel of deton 3680 m/sec at 1.02 (Marschall, v 2 (1917), p 493).

Decomposition Number of Hydrogen Peroxide in the case of the concentration of peroxide after being heated at 90°C for 24 hours to the original concentration (CIGS 30-115, p 9).

Decomposing Agent (Zerkleerungsmittel). According to Pic Area Tech Rep 1355 (1945), p 30 the following compositions were found in some German ammunition captured during WWI:

- a) Tin 60, lead 38, diamant 1.8 and nitromon 0.2%; used in some 37 mm HE shells.
- b) Tin 61 and lead 39%; used in some 40 mm HE shells.

Notes: According to E. Englehard, The Ordnance Sergeant, 1st of the 1st German Detachment agent consisted of a lead wire wrapped around the propellant bag of placed on top of it. (Upon detonation of the charge the wire formed a brittle alloy with the copper of the rotating band, and this alloy was rubbed off by the inner surface of the gun barrel. These particles then charge commencing, no decomposing agent was used. The shell shattered the brittle alloy, thus clearing the gun bore).

Delay Burning Process. See Zerkleerungsvorhaben.

Detonation Temperature Test (Versuchungs-Prob). See Ignition or Explosion Temperature Test.

Delay Compositions (Versuchungsschmelzen). A brief description of such compositions is given in the general section.

Shortly before WWI, the Germans developed gasless delay compositions suitable for electric detonators. These mixtures consisted of powdered potassium permanganate (KMnO₄) and saltpetre (Sb). Following is a brief description of the method of preparation as conducted at the Trier plant:

Procedure:

The dry crystalline K permanganate was ground in a special mill (called Kallmühle) to a particle size of about 0.006 mm. The saltpetre, received at the plant in a fairly finely divided state, was ground, without previous drying or other treatment, in a special mill (called Schwingmühle). The resulting powder was separated in an air classifier into fine (grit size under 40 microns) and coarser fractions. The coarser fraction was placed on a vibrating sieve containing 16,900 meshes per cm² and the fraction retained on the sieve was used as the coarse Sb. For the preparation of quick burning mixtures the fine Sb was used, while for slow mixtures the coarser material was more suitable. For instance a mixture of 36% fine Sb with 64% KMnO₄ ignited from 10 delay elements (v) burned 15.7 to 4.3 seconds, while the mixture of 36% coarse Sb and 64% KMnO₄ burned in 6.5 to 7.5 seconds. With a lower content of Sb and a higher content of KMnO₄ the burning time was longer. In order to obtain a composition with a desired delay, the coarse Sb was blended with the fine material.

Following is an example of the calculation for preparing a delay composition with a desired delay:

It is necessary to prepare 80 kg of delay composition consisting of 36% Sb and 64% KMnO₄ which would burn for 4.8 sec in a No. 10 delay element. The time of burning of coarse material is 7.50 sec and of the fine 3.50 sec.

If the "rectangle method" is used for composition (as is customary in Germany and some other countries of Europe) the calculation will be made by setting up the data above like:

7.50 1.35 2.65 4.85 (coarse Sb)
3.50 1.35 2.65 (fine Sb)
4.85 sec

In this configuration 1.35 is the difference between 4.85 and 3.50 and 2.65 is the difference between 7.50 and 4.85 seconds.

From the above, X may be calculated as follows:

$$X = \frac{1.35 \times (80-X)}{2.65} = \frac{1.35 \times 80 - 1.35X}{2.65} = \frac{108 - 1.35X}{2.65}$$

$$2.65X = 108 - 1.35X \text{ or } X = \frac{108}{2.65 + 1.35} = 27 \text{ kg (coarse Sb)}$$

The amount of fine material is then (80-X), or (80-27) = 53 kg.

After thoroughly mixing 27 kg of coarse Sb with 53 kg of fine Sb, a small sample consisting of 36 parts of mixed Sb and 64 pts of KMnO₄ was prepared and tested in a No. 10 delay element. If instead of the desired time of 4.85, 5.15 sec was actually obtained, then this Sb mixture would need to be corrected by adding some fine Sb (3.50 sec). The amount of fine Sb to be added was calculated with the "rectangle" method as described above and a small sample of new, corrected, mixture was prepared. If the burning time in a No. 10 delay element was exactly the desired 4.85 sec, the total batch consisting of 36% of "corrected" Sb and 64% of KMnO₄ was blended and pelleted. The pellets were ground and screened using sieves of 225 and 961 meshes per cm². The material which passed the 225 mesh sieve and was retained on the 961 mesh sieve was removed to storage while the material which was retained on the coarser sieve was ground and rescreened as above. The fine material (dust) which passed through the 961 mesh sieve was saved for addition to compositions considered to be too slow burning.

- a) Before commencing to load a delay element (v) with the above prep mixture, it was tested as follows:
 - i) Moisture content: A weighed sample of a delay mixture (5-10g) was heated for 2 hours at 110° in the loss of weight balance. The entire batch of delay composition was dried for several hours at 50° in a vacuum heated oven before it was loaded into delay elements.
 - ii) Particle size of Sb: A weighed sample of a delay mixture was loaded in a Gohmert-type sifter with bar water to remove the KMnO₄ and the particle size of the dried weighed Sb powder was determined (Kafe, 2 and 3).

Notes: The method for determination of particle size is not described in the reference given below.

A different type of delay composition consisting of K₂C₂O₄ (PbO₂) and silicon was used for the 200 mm HE mortar bomb. The composition in the mortar was: KC 3.5, red lead 75.5 and silicon 10.0%, while in the pellets it was: KC 2.7, red lead 72.0 and silicon 25.3% (Ref 1).

- a) R. M. Tomlinson Jr, Pic Area Tech Rep 1355 (1945), p 30
- b) R. A. Kerkhoff, B I O S Final Rep No 833, H M S O, London (1946), para 2, p 437 or 437/2
- c) Army, PB Rept No 95,613 (1947) (Manufactures of German Detonators and Detonating Compositions).

Delay Elements (Versuchungselemente). The elements used

during WWI consisted of metallic sleeves (of Al, Cu, brass, or coppered Fe) loaded with "gasless delay composition" (q v) consisting of powdered KMnO₄ 64 and Sb 36%. The sleeves had an inside diameter 3.30 to 3.45 mm, and an outside diameter of 6.45 ± 0.02 mm. The length (L) of the sleeves when using brass was as follows:

Delay in sec	1	2	3	4	5	6
L in mm	5	5.5	8	10.5	13	15.5
Delay in sec	7	8	9	10	11	12
L in mm	18.5	21.2	24.2	27	29.5	32

Loading of the sleeves was done by means of a 70 ton hydraulic press at pressures of 500 kg/cm². Details of the method are given in Ref 1, section 7.

The above delay elements were used in electric detonators, described briefly under: Detonators (Electric).

References:

- a) R. A. Kerkhoff, B I O S Final Rep No 833, H M S O, London (1946)
- b) Army, PB Rept No 95,613 (1947), Sections F & G.

Detonating Charge (Sprengladung oder Sprengkopf). The following charges were examined during WWI by U.S. Ordnance Dept establishments:

- a) Bohrspatron 28 (Blasting cartridge pattern 1928). A cartridge 3.5" long and 1.2" diameter, consisting of 3% oil of TNT or P.A. wrapped in varnished paper.
- b) Sprengpatrone, 28. A cartridge 4.1" long and 1.4" diam., consisting of P.A. wrapped in varnished paper.
- c) Sprengkopf 28 (Detonating block pattern 1928). A block 2.5" x 2" x 1.5" consisting of 2 lb of TNT or P.A. wrapped in varnished paper.
- d) Sprengkopf 28 consisting of two blocks of TNT, total wt 3 lb placed in a bakelite container ± 1.8 x 2.2 x 2.2.
- e) Sprengkugel 24 (Detonating block in container, pattern 1924). A block of TNT or P.A. weighing 2 lb 3 oz placed in a rigid container 7.9 x 2.9 x 2.2.
- f) Sprengkugel 24. A block of 90/10 PETN/Wax weighing 2 lb 3 oz.
- g) Gelblichladung 10 kg (Concentrated charge 3 kg). The detonating charge consisted of several blocks of TNT or P.A. with a total weight of 6.5 lb, placed in a zinc container (7.7 x 6.5 x 3") provided with carrying handle.
- h) Gelblichladung 10 kg. Same as above except that it contained 22 lb of TNT. The size of zinc container was 10 1/8 x 7 1/2 x 5 1/4.
- i) 12.3 kg Detonating Charge: A triangular block of 7 lb RDX/TNT in a stainless steel container.
- j) Platten. A block of plastic explosive RDX/Oil weighing 1 lb 15 oz.
- k) 300 g Hohlladung (Hollow charge). A shaped charge of a HE; size 3 1/2" high and 2.8" diameter.
- l) 400 g Hohlladung. A shaped charge consisting of 12 oz of PETN/Wax in an aluminum case 3 1/2" high and 2.8" in diam.
- m) 12.5 kg Hohlladung. A shaped charge consisting of 28 lb (with a container) of TNT in a sheet iron case 8 1/2" high and 1 1/2" diameter.
- n) 13 kg Hohlladung. A shaped charge consisting of 21 lb 3 oz (without a container) of 50/50 RDX/TNT in a mild steel container 8 1/2" high and 1 1/2" diameter.
- o) 50 kg Hohlladung. A shaped charge consisting of

110 lb (with a container) of TNT in a sheet iron case 10 1/2" high and 2" diameter, provided with a carrying handle.

- p) 500 g Hohlladung (Magnetic sensitive hollow charge). A shaped charge of a HE weighing 1 lb 1/4 oz of 3 lb Hohlladung, shaped charge consisting of 1 lb 50/50 RDX/TNT mixture in a metal container 7.7" high and 6.2" diameter.
- q) 3 1/2 kg Hohlladung. A shaped charge consisting of 2 1/4 lb of TNT in an aluminum container.

References:

- 1) Picatuney Arsenal Technical Rep No 1355 (1945), p 31
- 2) U.S. War Dept Technical Manual FM 5-25 (1945), PP 129-132
- 3) Dept of the Army Field Manual FM 5-25 (1954), pp 156-7.

Density of Fragments Test. See Fragments Density Test.

Durum Mining Association Testing Station. See under: Galleries, Testing, in the general section.

Detonationsdruck (Blast Pressure). See general section.

Detonationsfähigkeit (Ability or Detonant or Sensitivity to Initiation). The term is used to express the smallest numbered standard cap required to initiate the explosive under test. For instance, in Naudin's book Schieß- und Sprengstoffe, 1927 p 121, it is said that in order to initiate Ammonium ? or No 3 cap is required; while for Ammonium 1 and 5, 8, 10 cap suffices. This means that Ammonium 2 is less sensitive to initiation than are ammonium 1 and 5. The same test is used in Italy.

Detonationsgeschwindigkeit (Velocity of Detonation). See general section.

Detonationsübertragung Schwingung (Transmission or Detonation, Striking, etc.). Also called "Synthetic Detonation" (Artificial Detonation) (Ref 1) defines detonations (Detonations) as reinforced blasting caps which are designed to initiate explosives which are difficult to detonate by means of ordinary blasting caps.

The following military detonators were examined at Picatuney Arsenal during WWI and described in Ref 4, p 30:

Detonator R. contained 4 grains of 75/25 A/L/Sb: mixture over 6.9 grains PETN.

Detonator T. contained 3.9 grains of 42/58 L/A/L/Sb mixture over 10.8 grains of PETN in an Al cap. Both detonators were used in HE hand grenades.

Some of the captured German detonators in boxes (some times called gaiters) examined at Picatuney Arsenal during WWI are listed in Table 11.

Following are the principal current commercial detonators and blasting caps:

- a) Spherophat. A consists of an Al shell, 11 mm long, 4.35 mm in diameter, filled with a 6 mm layer of PETN weighing 0.11 g (0.04 lb) charge and a 3 mm layer, weighing 0.16 g of 70/30 (A/L/Sb) mixture, called in Germany the "Minutens" (Primer minuters) (Ref 1).
- b) Secondaries charges were pre-loaded at 450 kg/cm² (Ref 6).

Table 11
Detonators

Designation	Upper charge	Intermediate charge	Lower charge	Uses
Galus A	L.A. 52.5, S. 41%	RDX	RDX 92, was 8%	Not indicated
Galus B	L.A. 69.5, S. 31%	RDX	RDX 92, was 8%	"
Galus Model 40	L.A. / L.S.	RDX	PETN 87, was 13%	"
Detonator Galus	M.G.	Tonyl 49, TNT 51%	PETN	Lead line 37 mm HE and 30 mm HE shells 47 mm APEN shell
	L.A. 82.5, S. 7%			
	and similar 11%			
	L.A. With cover			
	charge of black powder			
	L.A. 144 and			
	L.S. 85.6%			
	L.A. 55, L.S. 45%			

Springshield, B. consists of an Al shell, 17 mm long, 7.9% in dia, filled with a 6 mm layer of PETN weighing 0.40 g (base charge) and a 4 mm layer weighing 0.40 g of "initiator" (primary charge) (Ref 6).

Notes: In both above caps the L.A. was of technical grade, containing 92-94% of PbO, and not more than 0.55% moisture.

Some of the "curved" (conical) caps are described in Ref 7. The so-called "Normal" copper cap No. 8 (Kupfer-Normalgeschosskopf No. 8) consists of a Cu shell, 6.3 mm in dia, press-loaded at 480 kg/cm² with 0.7% TNT (base charge), placed in two layers each weighing 0.35 g and with 0.55 g of MF as the primary charge. The same Ref 7 compares the properties of (flat-bottomed) caps with those of shaped charges. While the TNT test value and East crater test values are practically unaffected by a change in the shape of the bottom, the lead plane test values in each higher for the shaped charge.

Alus, (Ref 8) describes the following German detonators: Detonator B (Ref 8) consists of a shell 40 mm long, 6.3 mm in dia, filled with 0.6 g Tonyl compressed at 2000 kg/cm² (base charge) and 0.5 g of L.A./L.S. mixture (primary charge). Detonator No. 10 of D.A.G. Troisdorf contained 1.25 g of Tonyl and 0.5 g of L.A./L.S. mixture.

Abbreviations: L.A. Lead azide; L.S. Lead styphnate; M.F. Mercury fulminate; AP. Ammonium picrate; HE. High explosive; PETN. Pentaerythritol tetranitrate; RDX. Cyclotrim, or Hexogen; TNT. Trinitrophenol, References:

- 1) A. Sternbacher, Schiess- und Sprengstoffe, Leipzig (1933), pp 348-352.
- 2) C. Beyling & K. Dreckopf, Sprengstoffe und Zündmittel, Springer, Berlin (1936), p 151.
- 3) PB Repts 11,544 (1945), part III, p 10.
- 4) Dictionary Arsenal Tech Repts 155 (1945), pp 30-31.
- 5) A. Sternbacher, Spreng- und Schiessstoffe, Zürich (1946), p 105.
- 6) W. Schaefer, Sprengtechnik, No 10/11, p 186 (1932).
- 7) J. Kirsche, Sprengtechnik, No 12, pp 228-32 (1932).
- 8) Technical Report TM 9-1903-5 (1935), pp 547, 548, 566, 568, 569.

9) A. Lanza, Manuale del Maestro Esplosivista, Napoli, Milano (1935), p 77.
(See also BKOS 7 and Repts 644 and CIOB Rept 24-3).

Detonant (Detonative), A type of permissible explosive used before WWI. Some compositions are given in Table 12.

Table 12

Composition and some properties	Detonit 3 (powdered)	Detonit 5	Detonit 6 (w/Al)	Detonit 6 14
An mixture	82.7	64.0	82.0	82.0
K nitrate	-	-	-	10.0
NG (mixed with NC)	4.0	4.0	-	-
NG (neutral)	-	-	4.0	4.0
Aromatic nitro-compound	1.0	-	-	-
Vegetable meal	4.3	2.0	-	1.5
Wood meal	-	-	2.0	-
Coal (powdered)	-	4.0	0.5	-
MNN	-	-	1.0	2.5
Alkali chloride	-	22.0	-	-
No chloride	8.0	-	10.5	-
Oxybenz Balance	110.3%	-4.8%	110.9%	113.6%
Tuyal Test	225cc	220cc	230cc	235cc

Abbreviations: MNN. Mononitronaphthalene; NC. Nitrocellulose; NG. Nitroglycerin.

References:

- 1) J. Maass, Schiess- und Sprengstoffe (1927), p 146.
- 2) Maass, Nitroglycerin (1928), pp 434-5.
- 3) Beyling and Dreckopf, Sprengstoffe und Zündmittel (1936), p 141.

Diethyleneglycolinitrate (Diethyleneglycol Dinirate). See Diglykolinitrate.

Dinitro acid EDD (Ethylenediamine-Dinitrate). See general section. EDD was used by the Germans in Filler No 20, No 83, No 94 and No 86 as well as in the following mixtures of unknown name:

- a) EDD 45 and Am nitrate 55%
- b) EDD 45, Am nitrate 53.5% and Al 1.5%.

Notes: Mixture of EDD and Am nitrate forms a eutectic which permits cast loading.
References: Allied and Enemy Explosives, Aberdeen Proving Ground, Md (1946), p 145.

Dinitrobenzoperschlorot (Diazobenzoperschlorot). See general section.

Dinitrobenzomethylperschlorot oder Nitrobenzomethylperschlorot, known also as Blaupulver is described in the general section under Diazobenzoperschlorot.

Dinitro (Denarit). See general section.

Dicyandiamide (Dicyandiamide). Its manufacture in Germany is described in CIOB Final Report 1720 (1947). See also in the general section.

Didi-Pulver. An abbreviation for Diglykoldinitratpulver (Diethyleneglycolinitrate Propellant) (Sternbacher, Spreng- und Schiessstoffe (1946), p 44).

Diesel Ignitor. See Fuel Oil Ignitor.

Diethyleneglycolinitrate. See Diglykolinitrate.

Diethyleneglycol, Hexamine. See general section.

Diglykoldinitrat, "Diglykolinitrat" oder Didi (Diethyleneglycol Dinirate) (DEGN or DEGIN). Preparation and properties are given in the general section.

Following is a brief description of the German method of prep as practiced at the Künzels Fabrik of D.A.G.: 420 kg of technical "Diglykol" (DEG), consisting of 1% of diethyleneglycol and about 0.1% of water, was run slowly with stirring into 1218 kg of mixed acid consisting of 65% nitric acid and 35% sulfuric acid. The acid was cooled to below 25° by being circulated in cooling coils. Total time of nitration was 22 minutes.

Notes: A great excess of nitric acid was used in order to retard the decomposition of the otherwise extremely unstable spent acid. While the NG spent acid remained fairly stable for days, the DEGN acid had to be worked up at once since it decomposed rapidly on standing.

9) After the reaction was complete, the mixture was cooled to 15° and transferred to a separator where it was allowed to stand for 7 minutes. The spent acid (nitric acid 8-9, sulfuric acid 64-66 and nitrated product 4-7%) separated at the bottom, while the oil collected as the upper layer.

5) The spent acid was then transferred to a "desintegrator", while the oil, was run into the "primary washer" containing 100 liters of water stirred by air. The resulting acidic wash water contained an appreciable amount of nitric acid and was later desintegrated.

6) The oil was run into the "main washer" to be treated (with vigorous air-mixing) first with 500 l of cold water, then with 150 l of 5% soda ash solution, poached to 60° and finally with 1 M cold water.

7) A sample of the oil thus purified was sent to the laboratory and if the K test at 82° was not less than 20 min the material was considered to be satisfactory for use in the prep of the so-called Rohpulvermasse (q.v.).

The yield of DEGN was 710-715 kg or 170% of the DEG used; theoretically it should be 777 kg.

The purified DEGN had the following properties: light yellowish oil, d₄ 1.38 to 1.39, N content 14.1 to 14.2%, f.p. below -10°, v.p. (decomp ca 162° and puff at ca 200°), caloric value 1070 kcal/kg (vs 1715 for NG), water calculated as liquid, impact sensitivity with 2 kg weight 150 cm (vs 4 cm for NG), solubility in water ca 0.4% at room temperature, and volatility ca 4-5 rings more volatile than NG.

DEGN was used in the so-called "cool" propellants, such as "G" Pulver and "Gudol" Pulver, References:

- 1) O.W. Stickland, PB Rept No 925 (1945), p 57.
- 2) A. Sternbacher, Spreng- und Schiessstoffe (1946), pp 61-2. (See also CIOB Rept, 28-61).

Diethylenammonium Nitrate. See Di-Salz.

Diethyleneglycolinitramine (DMEDNA). Described in the general section. It was investigated by G.Römer, PBL Repts 85,160, p 14 as a component of some explosive compositions, such as:

- 1) DMEDNA 12, RDX 50, R-Salz 36, DPA 1 and uncrowned 1%
- 2) DMEDNA 2.5, RDX 96.5 and DPA 1.0%.

Diethylenetriamine (DMNA). Described in the general section. It was investigated by G.Römer, PBL Repts 85,160, p 13 as a possible addition to R-Salz in order to render it compatible at temps of 100° or lower; it was decided that incorporation of about 10% of DMNA was sufficient to give satisfactory results.

Dian. German abbreviation for Diazonaphthalene.

Dinitramin (Dinitramin) (DNA). Described in the general section under Asiline. The Germans used DNA during WW I as an additive to TNT. The resulting explosive was yellow in color, less powerful than TNT, and much less sensitive to impact or friction. It produced larger projectile fragments than did TNT (Allied and Enemy Explosives, Aberdeen Proving Ground (1946), p 93).

Dinitraminol oder Dial (Dinitronitrolole) (DNAs). See general section under Anisole; was used by the Germans in some explosive compositions, such as "Anisole No 40" (q.v.).

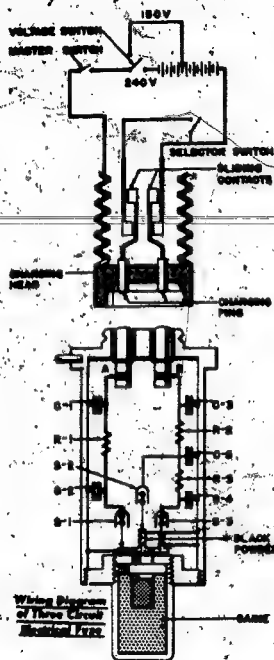
Dinitrobenzol (Dinitrobenzene) (DNB). See general section under Benzenol. It was used by the Germans as an extender for TNT and as a substituent in some explosives, such as RDX. The addition of it to some high-melting explosives rendered them suitable for cast loading (Allied & Enemy Explosives, Aberdeen Proving Ground (1946), p 111).

Dinitroethylol. See Diglykoldinitrat.

Dinitroethylenhydrazin (Dinitroethylenhydrazin) (DNCH or DNCHG) is described in the general section under Chloroethylen.

Dinitroethylol (Dinitroethylol). See general section, under Glycol.

Dinitronaphthalin, Dian. (Dinitronaphthalen) (DNN). See (q.v.).



If the selector switch was held open, then this charge was through plunger B to the storage condenser C-3 and nothing passed to the instantaneous circuit. The circuit passed through the resistance R-2 to the condenser C-3 because of the delay train through both resistances R-2 and R-3 to firing condenser C-4. If the bomb had been dropped from an altitude of less than 1170 ft, the latter circuit would not be closed before impact and the igniter bridge associated with the trembler switch S-2 would fire the long delay pellet which actuated through the explosive train of the fuse would detonate the bomb. If the bomb was dropped from an altitude greater than 1170 ft, high circuit would be armed before impact, but because of the character

delay train used in conjunction with the trembler switch S-3, the short delay would initiate the final explosive train. Electrical time fuse (EIZZE), contained essentially the same basic parts as the electrical impact fuse (EIAZ), except that the trembler switches were replaced by a vacuum tube which became conducting at a critical predetermined voltage. At the instant the bomb was started on its trajectory, an electric charge was put on the storage condenser, and another smaller charge was put on the firing condenser. The time setting of the fuse was adjusted by varying the amount of charge placed on the firing condenser. During flight, part of the charge on the storage condenser leaked through the resistor to the firing condenser. As the charge on the firing condenser increased, the voltage across the vacuum tube also increased. When the firing voltage of the tube had been reached, the firing condenser discharged through the tube and the igniter bridge thus firing the fuse.

Electrical bomb fuses are described in Refs 1 and 3, and are listed in this work under Fuse. Some of these fuses are described in this work under Aerial Bomb Fuse. An electrical time fuse (EIZZE 5/30) for use in projectiles is briefly described in Ref 4, p. 605-B. Prior to firing the projectile, the fuse was charged either by hand or by a machine by putting 300 to 500 volts across the shell and an insulated contact which put voltage on the storage condenser. The charging could also be done by allowing the "feeler wire" (connected to the electrical circuit of the fuse) to connect the "muzzle charging ring" or the projectile was leaving the gun. A brief description of a muzzle charging ring is given in Ref 4, p. 606.

A device, described in Refs 2, 422 and 4, p. 623 as the electric fuse, ERZ 35, was used for igniting the black powder charges which set off the propellant of 15 cm and 21 cm rockets. This device is briefly described in this work under Rocket Propellant Igniter.

(See also under Electrical Igniter and under Igniter).

References:
1) Army War Dept Tech Manual TM 9-1983 (1942), Heavy Bombs and Fuses, File Numbers 2321-5, 2321-6, 2324-92, 2324-93, 2324-94.

2) Army Ordnance Bomb Disposal Center, Aberdeen Proving Ground, Md (Nomenclature: German Artillery Projectiles and Fuses p. 42).

3) Army War Dept of the Army Tech Manual TM 9-1983-2 (1953), German Bombs, Fuses, Rockets, etc., pp. 125-132 and others.

4) Army War Dept of the Army Tech Manual TM 9-1983-5 (1953), German Projectiles and Fuses, pp. 605-7 and 623.

Electric Fuse Primer Composition. See Primary and Initiating Compositions.

Electric Igniter (Elektrischer Zünder). Among the numerous igniters used by the Germans in mines was one type, EMEZ 35, which used an electric circuit for setting the fuse to a mine. This fuse is briefly described in TM 9-1983-2 (1953), p. 300-1.

(See also under Igniter).

Electric Igniter and Primer (Elektrischer Zünder) Used for Commercial Explosives. These devices, described in Beyhag-Drehtopf, Sprengstoffe und Zündmittel (1936) may be subdivided into the following groups:

- Einfache Zünder (Simple igniter). It consisted of a capsule (Hülse), a priming composition (Zündsatz) and electric lead wires connected to a bridge wire (B R'D), pp. 177-172.
- Zusammengesetzter Zünder (Composite igniter or primer), such as Sprengzünder (detonating primer), consists of a simple electric igniter combined with a detonator, (B R'D), pp. 174 and 221-24.

c) Zünder mit fest eingeregelter Sprengkapsel consists of a simple primer into which a No 8 blasting cap (Sprengkapsel No 8) is firmly set (See B R'D, pp. 174 and 225).

d) Unterwasserzünder (Underwater primer) is described in B R'D, pp. 225-26.

Zeündschützvorrichtung (Time igniter with fuse) consists of a simple primer combined with at least a 20-cm piece of fuse (B R'D, pp. 175 and 226-29).

f) Schweißzünder (Inexplosive igniter or primer), described in B R'D, pp. 17 and 225.

g) Unterwasser-Schweißzünder (Underwater instantaneous igniter or primer), described in B R'D, pp. 175 and 225.

Abkürzung: B R'D Beyhag and Drehtopf.

Electric Matchless or Flashless is the combination of bridge wire, igniter head and lead-in wires employed in electric blasting caps and detonators.

(CLOS Sept 24-3, p. 7) Also under F. Schenck Manufacturing.

Electric Proximity Fuse. See Proximity Fuse.

"Eltion" (Elephant). A tank destroyer known also as Schärer Panzer Jagd "Eltion". It was an improved version of "Ferdinand" (q) - See also under Panzer.

Elektronenbombe (Electron-bomb). See general section.

Empfindlichkeit gegen Reibung (Sensitiveness to Friction). See general section.

Empfindlichkeit gegen mechanische Einwirkungen (Sensitiveness to Mechanical Action). See general section.

Empfindlichkeit gegen Stoss (Sensitiveness to Shock or Impact). See general section.

Empfindlichkeit gegen Wärme (Sensitiveness to Heat), also called: Chemische Stabilität (Chemical Stability) is described in the general section under Stability.

Entzündmittel für nachgelassen Pulver. See Entry Content of a Propellant Charge.

Engrit (Engrit). According to Neeson (Ref 1), Engrit was a commercial explosive made after W. H. Nobel's Dynamic A-G. The explosive was prepred by wet grinding various kinds of surplus double-base propellants in "Explosive" mills between steel discs, to a particle-size of 0.5 to 1 mm, followed by drying and packing in cartridges 2 1/2 to 3 1/2 mm diameter. This explosive was used to a great extent in parashooting.

According to Pepin Lebelleur (Ref 2), Engrit and Trielwastel were industrial explosives prepared by blending a smokeless propellant (left as surplus after W. H. Nobel's previous work) with about an equal quantity of a sulfur such as sulfur or acetone, with liquid aromatic nitro-compounds and oxidizing agents such as alkali nitrates or chlorates in a blender. The strength of these explosives as determined by the Traut test was 330 to 350 cc. velocity of detonation 3000 to 5000 m/sec.

References:
1) P. Neeson, Nitroglycerin, etc., Baltimore (1928), p. 449.

2) J. Pepin Lebelleur, Powder, etc., Paris (1955), p. 437.

See also Nitroglycerin-chemical Explosives (Mining Lists 33, 35 and 36) and also Trielwastel, SN.

Entry Content of a Propellant Charge. According to PB 925 (1945), p. 82, the energy content is equal to the charge weight of a propellant multiplied by its caloric value. For a given propellant and a given initial (muzzle) velocity, the energy content is constant and independent of the type of propellant used. For instance, if for a certain initial velocity of a projectile the charge weight of a propellant with a caloric value of 820 cal/g is 43 g, a propellant with a caloric value of 570 cal/g (such as a nitroguanidinium) would require a charge of 62.4 g. (See Effectivity-Caloric Value of Propellants).

Entzündungspunkt (Flash Test). The test is applied to smokeless propellants is described by H. B. Bannan, Das Schießpulver, (1926), p. 304.

Entzündungspunkt oder Entzündungstemperatur (Flash Point, Kindling Temperature). The test is described in the general section.

Entzündungsmittel. See Detonating Agent.

Entzündungsmittel (Antiflash Type Igniter with HE Charge). See under Igniter.

Entzündung oder Trocknung (Dehydration, Drying). See general section.

Enzlin Rakete (Enzlin Rocket). One of the guided rockets developed and used by the Germans during WW II, it has been described by:

1) F. Röss, J. Guided Missiles, Rockets and Torpedoes, NY (1946), p. 43.

2) A. Durosoy, Les Armes Secrètes Allemandes, Paris (1947), p. 99.

3) TM 9-1985-2, p. 229-32.

(See also Great Enzlin or E-4 Missile).

Entzündlichkeit (Inflammability). See general section.

Entzündungsmittel (Ignition Mixture). See general section.

Entzündungspunkt (Ignition Point). See general section.

Entzündungspunkt (Ignition or Burning Point). See general section.

Entzündungstemperatur oder Vapourisationstemperatur (Ignition, Deflagration or Explosion Temperature). See general section.

Entzute (Earth Quake). A rock-resistant 1800 lb armor-piercing bomb (PC 1800 RB), used by Stuka bombers against land targets. This bomb is mentioned, but not described, in TM 9-1985 (1942), File No 2324-92.

Erosionless Priming and Initiation (Erosionfreie Zündung). Priming and initiating compositions containing mercuric fulminate and the chlorates (such as KClO₃) have been known to cause considerable erosion of gun barrels. In 1904, H. Ziegler of Switzerland, therefore, proposed that the salts such as the nitrate be substituted for the chlorate salts. These new compositions were known in the industry as "positive Zündungen" (positive primers). As these substances were not entirely satisfactory, further search resulted about 1930 in the invention of compositions based entirely on organic compounds, such as, Tetrazene (Tetrazene). These substances, called "erosionfreie Synchronzündern" were manufactured before WW II by the Rheinisch-Westfälische Sprengstoffe A-G in Germany.

a) Flam C 250 A (B or C) contained 50 kg of oil incendiary mixture and TNT bursting charge (p 52)

c) Flam C 500 contained the incendiary oil consisting of 70% petroleum and 30% TNT, with TNT burning charge. (p. 54)

(See also Incendiary Bombs, Brandbommen and Sprengbommes). (Illustrations are given under Bombe).

Flammability Test (Entzündlichkeitsprobe). A special apparatus called "Flammenpendel" and its application to testing of various explosives and pyrotechnic compositions was described by P. Leese *SS* 27, 366-68 (1932).

Flammgeschwächungsmittel (Flame Extinguishing Additive, or Flame Reductant). See Flame Reducing Compounds in the general section.

Flamethrower (Flange Projectile), called also "Square-bore", or "Littajoko" was a sub-caliber projectile provided with a flange and three hollow studs as shown on Fig. 1. It was described in the TW 9-1981-3, p. 360.

It was fired from a cylindrical rifled barrel to which a smooth-bored, tapered muzzle extension was attached.

The principal advantage of the "flange" projectile in comparison to the other subcaliber projectiles was that it had no parts to be discarded, because the hollow stud and the flange were easily depressed when the projectile passed from the gilled section of the gun to the smaller caliber smooth bore extension.

(Compare with Arrowhead Projectile, Arrow or Needle Projectile, Disintegrating Band Projectile, Röchling Projectile, Sabot Projectile and Tapered Base Projectile)

Flare (Leuchtkugel oder Fackel). A German flare usually consisted of a cylindrical container housing an illuminating element. Upon being ignited by a pull friction igniter or a time fuse the flare burned vigorously producing intense light and heat. The illuminating element consisted either of a single or a multiple candle unit which varied in intensity of illumination and color. Flares were made with or without parachutes.

A brief description of the following flares is given in TM 9-1985-2 (1953), pp 65-81:

1) LC 10 (Leuchtzylinder 10) consisted of an aluminum cylinder, a single candle in a cardboard box, and 100

cylinder, a single candle in a carbide flare, an oxygen cylinder work fuse and a parachute located in the tail end. The flare was dropped from a plane and at a predetermined time the fuse fired and ejected the candle and its parachute from the body. Simultaneously the candle was ignited (p. 65)

2) FB 50, Single Candle Parachute Flare (p. 66)

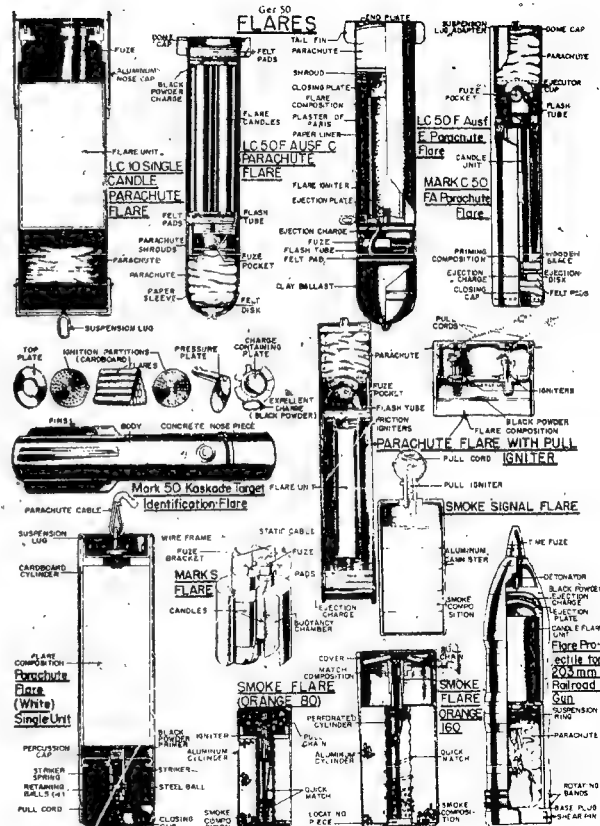
3) LC 50F Ausf. C, *Parachute Flare* consisted of a cylindrical body with a 100 mm diameter and 150 mm length attached by means of brass screws. On releasing the flare, the pyrotechnic delay (inside the fuse) was ignited. This fired the quickmatch, which in turn burned through the flash tube and ignited the black powder. The black powder was contained in a 100 mm diameter tube by the deflagrating black powder, caused all four flare candles and the parachute to be expelled through the nose, after shearing the holding screws. Simultaneously, the candles were ignited through perforations in the nose. The burning time of the candles was 7.5 sec. The nitrate 75.8 (Al 16.5 and 5.77%). The burning time was slightly over 3 min and the candlepower 216,000 (p 68).

4) LC 50F Ausf. E, *Single Candle Parachute Flare* consisted of a cylindrical body with a 100 mm diameter and 150 mm length attached by means of brass screws. On releasing the flare, the pyrotechnic delay (inside the fuse) was ignited. This fired the quickmatch, which in turn burned through the flash tube and ignited the black powder. The black powder was contained in a 100 mm diameter tube by the deflagrating black powder, caused all four flare candles and the parachute to be expelled through the nose, after shearing the holding screws. Simultaneously, the candles were ignited through perforations in the nose. The burning time of the candles was 7.5 sec. The nitrate 75.8 (Al 16.5 and 5.77%). The burning time was slightly over 3 min and the candlepower 216,000 (p 68).

5) LC 30F- Ausf G, Single Candle 'Parachute Flame'
(pp 69-70)

6) Mark C 50 F/A Parachute Flare consisted of a cylindrical aluminum housing containing a parachute, fuse, quicmatch, single candle unit, flash tube, priming composition and ejection disk. When the flare was released, the serial burst fuse started to function. The flash ignited the quicmatch and the flame was transmitted through the flash tube to the tail and to ignite the ejection disk of black powder. The pressure of the gases developed by the burning powder expelled the parachute and the candle through the nose. Simultaneously the primer composition and the candle were

7) Mark 30 Kushode Target Indicating Flare consisted of a sheet metal cylindrical container 7.7" diam and 41.0" long containing 62 flares (in three layers separated by perforated cardboard partitions), an expelling charge (a 100% pure potassium perchlorate propellant disk) and an igniter (fuse) assembly. A heavy concrete nose was provided to make the missile fall with the nose downwards, when released from a plane. As the missile fell, the expelling charge was ignited thus ejecting the flares (candles). At the same time the propellant



usually loaded in small bags separately from the propella.

Flash Reduction in Projectiles. When it was required by the German High Command to have an AA (Flak) projectile whose explosive flash is practically invisible in the night

...and the

Components and properties	Designation		
	Fordix 2	Fordix 3	Fordix 4
Amplitude	41.0	37.0	38.0

Am. nitrate	41.0	37.0	38.0
NG (nitroglycerin)	25.0	25.5	21.0
Colloid cotton	1.0	1.5	1.0
MNT (mononitrotoluene)	3.5	5.0	5.0

Glyceria	0.7	3.0	3.0
Cereal or potato flour	-	-	12.0
K chloride	22.0	24.0	19.0
Am oxalate	-	-	1.0
Bolin (china clay)	0.1	-	-
Dextrin	0.7	4.0	-
Oxygen Balance, %	-	-	-19.5
Truxal Test value, cc	-	-	220

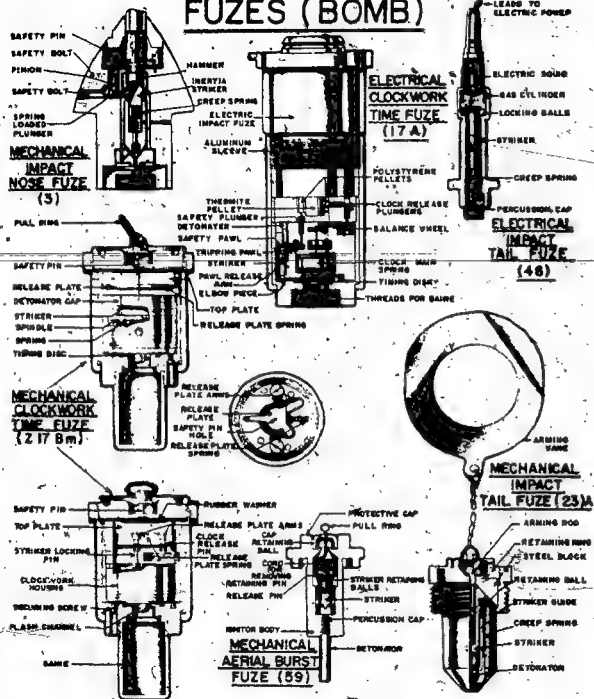
FB Rept No 78,271 (1947), p 4; 2) A.J. Roux. *Mém Pou*
34, 132 (1952).

Following are some values for the average distance to obtain one penetration per square meter using a 105 mm shell:

walled projectiles which being lighter than water float on its surface [Daniel, Dictionnaire (1902), p. 310].

Fritsche Zündschnur (Fritsche's Fuse). A core consisting of a pressed mixture of K nitrate 63, alderwood charcoal (Erlenholzkohle) 13, and pulverized sulfur 24% enclosed in a fabric tube. It was slow-burning. {A. Stur-bacher, Spreng- und Schussstoffe, Zürich (1948), p. 107}.

FUZES (BOMB)



File No 2321.5)

- 3) Elec Short Time Aerial Burst Fu EIAZ2 (9) or (9)* used in parachute flares and photoflash and gas bombs (Ref 4, p 167)
- 4) Elec Imp Fu EIAZ (35) or EIAZ C 50 (15) (obsolete) was used in SC 50 to 2500 kg, SD 50 to 1400 kg and SB 50 kg bombs (Ref 2, file 2321 and Ref 4, p 139)
- 5) Elec Mech Long Delay Time Fu EIAZ (17), Type 7, used in SC 250 and 500 kg bombs having two pockets (Ref 4, p 252)
- 6) Elec Mech Time Fu EIAZ (17A), EIAZ (17A)*, EIAZ (17B)* used in the same bombs as EIAZ (17) (Ref 4, p 254)
- 7) Mech Time Fu Z 172m used in SC 500 to 1000 kg, PC 1000 kg and BSB 1000 kg bombs and the 293 flying bomb (Ref 4, p 155)
- 8) Mech Imp Tail Fu (23A) used in Brand 10 kg, NB 2 kg and SC 3 kg bombs as well as in single unit parachute flares (Ref 4, p 134)
- 9) Mech Imp and Antishock Fu (24) and (24A) used in the forward pocket of SC 2500 bomb (Ref 4, p 135-8) (See a brief description under Antishock Fuze)
- 10) Elec Imp Fu EIAZ (25), (25A), (25A)* and (25A)** used in HE bombs (Ref 4, p 140)
- 11) Elec Imp Fu EIAZ (25)B, 25 B, (25) C and (25) D used in SC 50 to 500 kg and some for bombs (Ref 4, p 141-2)
- 12) Elec Proximity or Imp Fu, Special EIAZ (26) used in HE bombs (Ref 4, p 144)
- 13) Elec Imp Fu EIAZ (28)A, used in HE bombs SC 50 to 2500 kg and in SC bomb C 250 (Ref 4, p 163)
- 14) Elec Imp Fu EIAZ 28 (*) or EIAZ C 50 28 (*) used in HE bombs (Ref 4, p 163)
- 15) Elec Imp Fu EIAZ (28)B used in SC bombs against sea targets (Ref 4, p 163)
- 16) Elec Imp Fu EIAZ (28)B*, (28)B* and (28)B** used in HE bombs (Ref 4, p 163-4)
- 17) Mech Aerial Burst Fu (29) used in LC 101 parachute flare (Ref 4, p 164)
- 18) Elec Imp Fu EIAZ (35) used in HE and AP bombs (Ref 4, p 142)
- 19) Elec Imp Fu EIAZ (38), (38)mg and (38) used in HE bombs (Ref 4, p 165-6)
- 20) Elec Imp Fu EIAZ (38A) used in SC 250 kg bombs when employed on depth charges against U-boats. (Ref 4, p 166)
- 21) Elec Imp Fu EIAZ (38)B & (38)C used in FX 1400 and HE bombs (Ref 4, p 166-7)
- 22) Mech Antishock Device Zuezl 40, Typen 1, II & III used in SC 250 to 500 kg bombs under fuses (17A) or (17B) (Ref 4, p 157-8) (See a brief description under Antishock Fuze)
- 23) Mech Imp Fu AZ 41 or 34-41 used in SD 1A "Butterfly" bomb (Ref 4, p 132)
- 24) Mech Imp or Aerial Burst Fu AZ (41)A cor* was used in SD 2 B "Butterfly" bomb (Ref 4, p 132)
- 25) Elec Imp Fu EIAZ (45) used in unknown (Ref 4, p 142)
- 26) Elec Imp Fu EIAZ (45A) used in SC 50 bombs (Ref 4, p 142)
- 27) Electrically Armed Mech Imp Tail Fu AZ (46), used in KC 30 gas bombs (Ref 4, p 145)
- 28) Rocket Bomb Fu Assemblies (49)A & (49)B, Type 9 used in PC 2000, 1800 kg bombs and 1800 kg "Erdstark" (Ref 2, file 2324.92 & 4, p 169)
- 29) Rocket Bomb Fu Assembly (49)C used in PC 1800KS (Ref 4, p 170)
- 30) Elec Antishock Fu 50 and (50) used in SC 250 and 500 kg bombs in conjunction with fuses (17), (17)A or (17)B (Ref 4, p 181-3)
- 31) Elec Antishock Fu 56 q Y* (See under Antishock Fuze) used in HE bombs alone, or in conjunction with other Rheinmetall fuses (Ref 4, p 184)
- 32) Elec Imp Fu EIAZ C50 (5) (obsolete) & C/50 (15) used in HE bombs (Ref 4, p 197)
- 33) Elec Imp Fu EIAZ (55A)mp, (55A)M & (55A)* used in SL & SD and other bombs requiring instantaneous

Get 56

action (Ref 4, p 15-4)

- 34) Elec Chemical Time Fu EIAZ (57) used in "Sasha" bombs (Ref 4, p 157)
- 35) Mech Aerial Burst Fu (59) used in single & four candle parachute flares and BIC 50 photoflash bombs (Ref 4, p 171)
- 36) Elec Aerial Burst Fu 59A & (59A)* used in A/P and inc containers (Ref 4, p 172)
- 37) Elec Aerial Burst Fu (59)B used in some HE bombs and parachute flares (Ref 4, p 172)
- 38) Mech Aerial Burst Fu Z 56 used in supply-dropping containers (Ref 4, p 186)
- 39) Special Imp Fu 266 used in SD 10A bomb (Ref 4, p 146)
- 40) Mech Time Fu Z 677 Zeit used in SD 2B "Butterfly" bomb. It was located centrally in the upper longitudinal surface of the bomb (Ref 4, p 159)
- 41) Mech Time Fu (70) used in Mk AB "C" containers to ignite 2 of the 3 candle units housed in the container (Ref 4, p 160)
- 42) 2500 Aerial Burst Fu, Pyrotechnic Delay 69C II, 69D & 69E (Ref 4, p 173) used in AB 36, 500, 1000 and BDC 10 containers
- 43) Chem Mech Long Delay and Antishock Fuze (70A) used in SD 2B bomb (Ref 4, p 187)
- 44) Mech Antishock Fuze (70)B & (70)B/I used in SD 2B bomb (Ref 4, p 187)
- 45) Modified Mech Antishock Fuze (70)B used in aircraft towed parachute bomb (Ref 4, p 188)
- 46) Elec Aerial Burst Fu, Pyrotechnic Delay EIAZ 79, 79A, & (79A)* used in parachute flares, SC 250 & 500 bombs, A/P inc containers and photoflash bombs (Ref 4, p 174)
- 47) Mech Imp "All-Weys" Action Fu VZ (80) used in Ha 293 flying bomb (Ref 4, p 199)
- 48) Disso VZ (80A) used in V-1 flying bomb (Ref 4, p 190)
- 49) Mech Aerial Burst Fu Z (89) used in photoflash bomb, parachute flares and area containers (Ref 4, p 175)
- 50) Disso Z (89)B, (89)C & (89)D used in some containers (Ref 4, p 177)
- 51) Elec Imp Fu EIAZ (106)* used in Flying Bomb "Penne" missile 16 (Ref 4, p 146)
- 52) "Dust Fuze" used in SD 10 bombs (Ref 4, p 191) (See description under Di.)

Following are abbreviations and designations used for bomb fuses:

AZ	Aufschlagzünder	Impact fuze
EIAZ	Elektrischer Zeit-zünder	Electronic time fuze
EIZ	Elektrischer Zünder	Electronic (fuse)
LZ	Langzeit	Long time delay
VZ	Vorwarnzünder	Safety fuze
Z	Zünder	Fuze
ZLZ	Zeitzünder	Time fuze
Zu	Zusatz	Addition
ZLZ-St	Zeit-zünder-Strich	Fuze extension cap

Other German abbreviations are given at the end of this German section, following the Vocabulary

Several of the German bomb fuses were examined at Picatinny Arsenal as can be seen from the following report: A. A. Schilling, Pic Tech Rept 1572 (1945) (Chemical Long Delay Bomb Fuze, EIAZ)

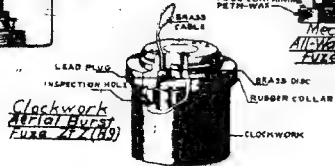
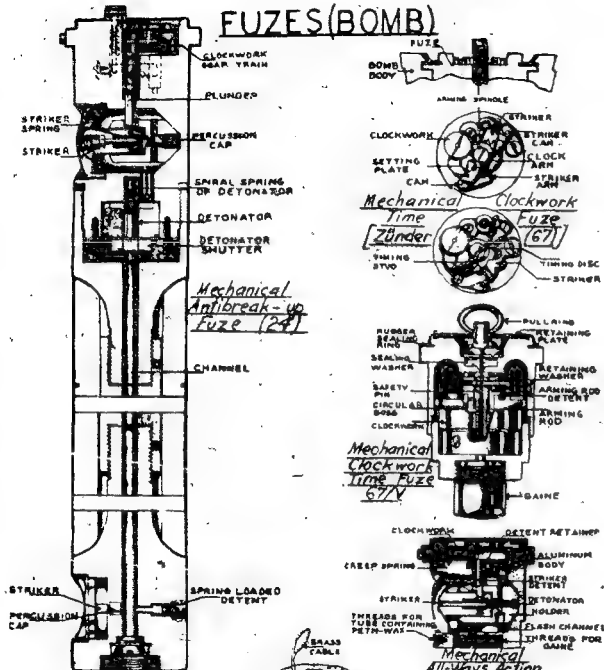
A. B. Schilling, ibid, 1574 (1945) (Mechanical Time Long Delay Bomb Fuze, L Zt 2)

C. A. Schilling, ibid, 1581 (1945) (Rheinmetall and Long Delay Bomb Fuze, EIAZ 35A)

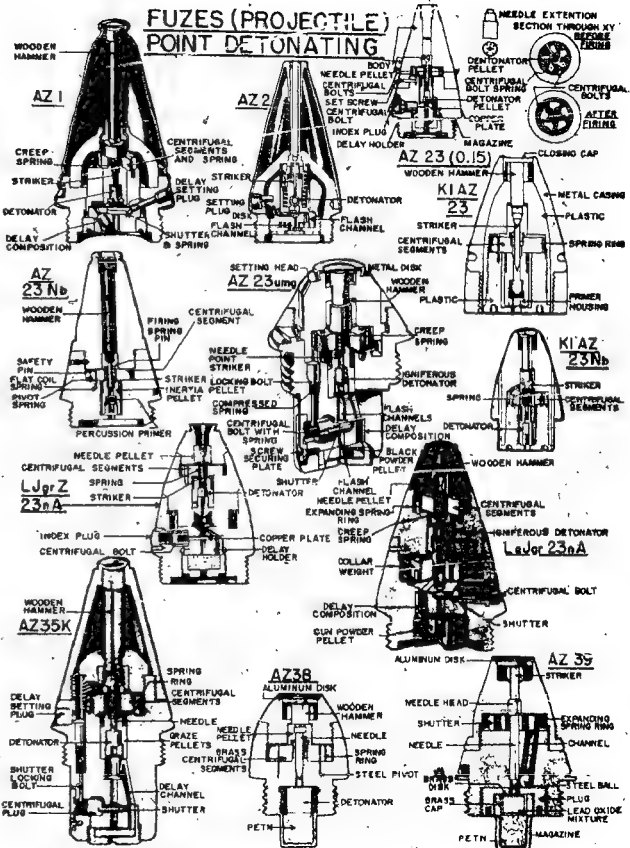
(See also Aerial Burst, Antishock and Electric Fuses)

B. Projectile Fuze (Geschosszünder) existed even in a greater variety than bomb fuses. The former may be subdivided into Point Detonating (P.D.) and Base Detonating (B.D.) types. A brief description of typical

FUZES (BOMB)



FUZES (PROJECTILE) POINT DETONATING



German projectile fuses is given by Engleburg (Ref 2). The following types are listed and briefly described in Refs 1 and 5.

1. Point Detonating Fuse

1. Imp Fa AZ 1 used in 75 mm and larger caliber shells (Ref 1, p 580).
2. Imp Fa AZ 2 used not indicated (Ref 1, p 580).
3. Perc Fa AZ 23 Series were the most important and used throughout for German Artillery Ammunition, mostly for 75 mm and larger calibers. All the different fuses bearing the number 23 were similar in functioning and major differences.

23 type fuses existed in the following variations:

- a) Perc Fa (with delay 0.15 and 0.25 sec) aluminum body AZ 23V0.13 and 23V0.23 used in shells for 75 mm Gun and 105 mm Howitzer (Ref 3, p 339 & 5, p 571).

- b) Perc Fa AZ 23Gub used in the 75 mm Mountain Gun (Ref 1, p 570).
- c) Perc Fa plastic body AZ 23V(0.15)P0 and AZ 23V0.23P0; used not indicated (Ref 5, p 533).

d) Perc Fa plastic body AZ 23N0P0; used in 150 mm Smoke shells (Ref 5, p 607).

- e) Perc Fa zinc body AZ 23V0.13X26 and AZ 23V(0.15)X26; used not indicated (Ref 5, p 573).
- f) Modified Perc Fa AZ 23ung used in 75 mm and 105 mm HE shells (Ref 5, p 575).

g) Perc Fa AZ 23/28 used in 88 mm HE AA shells (Ref 5, p 565).

- h) Perc Fa (delay 0.15 sec) AZ 23/42V(0.15); used not indicated (Ref 5, p 573).

i) Perc Fa (modified) 1/19Z 23A used in 75 mm Light Infantry Gun (another model of 1/19Z 23A was used in 210 mm Rocket 42 21 cm Vgr 42 Sp0 (Ref 1, p 583)).

j) Perc Fa and Perc Fa (delay 0.4 sec) 1/19Z 23 and 23V(0.4); used not indicated (Ref 5, p 566 & 5, p 573).

k) Perc Fa 1/19Z 23NB (also 23NB) used in Smoke shells (Ref 5, p 575).

Notes: Other, less important, versions of fuse 23 included: AZ 23 (obsolete), AZ 23V(0.15), AZ 23(0.2) and AZ 23(0.15)ung (Ref 5, pp 573-4).

4) Small Perc Fa AZ 18 Series existed in the following variations:

- a) Perc Fa (small) K1A2 23 used in 75 mm HE and 75 mm 105 mm Smoke shells (Ref 5, p 570).

b) Perc Fa K1A2 23NB used in Smoke shells (Ref 5, p 578).

- c) Perc Fa (with delay 0.2 sec, modified K1A2 23V (0.2)ung) used in 75 mm A/T Guns 40, 42, 76.2 mm Russian A/T Gun 36 and Field Gun 39 (Ref 5, p 574).

Notes: Other, less important, versions of small fuse 23 included: K1A2 23V(0.2), K1A2 23V(0.15), K1A2 23V(0.3) (P0) and K1A2 23N0P0 (Ref 5, pp 574 & 578).

5) Impulsive DA and Graze Type Fa (with a combined gas and DA mechanism) AZ 33K used in 170 mm HE Shell (Ref 5, p 580).

- 6) Mech Imp Fa AZ 33 used in HOC projectiles (Ref 5, p 532 & 5, p 568).

7) Detonating Imp Type Fa (with DA mechanism) AZ 39 used in 20 mm HE shell (Ref 5, p 337 & 5, p 569).

8) Imp Fa K1A2 40NB & 40NB(P) used in Smoke projectiles (Ref 5, p 570).

- 9) Perc Fa AZ 47 & AZ 48, similar in construction to AZ 19, were used in 20 mm Ammo (Ref 5, p 571).

10) Perc Fa AZ 49 used in 20 mm Shell (Ref 5, p 571).

11) DA Imp Fa AZ 150 & 150R3 used in 20 mm Shell (Ref 1, p 315 & 5, p 564).

12) Imp Fa AZ 150Z used in 20 mm Shell (Ref 3, p 303 & 5, p 547).

13) Imp Fa AZ 1503 used in 20 mm Shell (Ref 3, p 309 & 5, p 547).

14) Imp Fa AZ 1504 used in 20 mm Shell (Ref 3, p 309 & 5, p 547).

15) Imp Fa AZ 1531 used in 20 mm Shell (Ref 3, p 315 & 5, p 549).

16) Imp Fa AZ 1552 used in 15 mm Projectile (Ref 5, p 550).

17) Imp Fa AZ 1551 used in 15 mm Projectile (Ref 2, p 316 & 5, p 550).

18) Imp Fa AZ 1552 used in 15 mm Projectile (Ref 5, p 550).

19) DA and Graze Fa AZ 5045 used not indicated (Ref 5, p 552).

20) Imp DA Fa AZ 5045 used in 20 mm Shell (Ref 5, p 552).

21) Mech Imp Fa AZ 5072 used in 20/20 mm and 42/20 mm HE shells for Targeted Bore guns (Ref 5, p 315 & 5, p 553).

22) Imp Fa AZ 5075, AZ 5075 mk & DAAZ 5075 used in 37 mm Rodded A/T Bomb (37 cm Pak Scigun) (Ref 1, p 319 & 5, pp 554-5).

23) Imp Fa AZ 5095 used in 88 mm A/T HOC Rocket (Ref 5, p 555).

24) Imp Fa AZ 15H9 used in 150 mm Shell with BC (Ref 5, p 586).

25) Mech Time and Imp Fa Dopp Z 28K used in 210 & 180 mm projectiles (21 cm KGr 38 & 28 cm G-39) (Ref 5, p 605).

26) Mech. Time and/or Imp Fa Dopp Z 5/60 FI used in 88 mm and 105 mm HE AA shells (Ref 5, p 383 & 5, p 603).

27) Ditt0 Dopp Z 5/60; used not indicated (Ref 5, p 318).

28) Mech Time and Graze Action Fa Dopp Z 5/60G45 used in 170 mm Gun in Mortar Mounting (17 cm K 1 Mehl) (Ref 5, p 601).

29) Combination Fa Dopp Z 5/160Gub used in shells for Mountain Guns (Ref 5, p 596).

30) Supercompressive Imp Fa EKZ C/28 used in shells for Dattel guns (Ref 5, p 605).

31) Elec Time Fa ELIZAZ 5/30; used not indicated (Ref 5, p 601).

32) Imp Instantaneous and Delay Fa under BC HgZ 135D used in 210 mm Rocket (21 cm Vgr 42 Sp0) (Ref 5, p 585).

33) Ditt0 HgZ 135K used in 170 mm HE Shell (Ref 5, p 593).

34) Imp Fa (Russian Design) KTM-1 used in 76.2 mm HE Shell (Ref 5, p 577).

35) DA Detonating Type Fa KZ 14.7 cm Pak Sprg used in 47 mm HE Shell (Ref 5, p 565).

36) Mech Imp Fa (with a self-destructing arrangement) KZ 20PWP used in 37 mm HE AA Shell (Ref 5, p 557).

37) DA Mech Imp Fa (with a safety device which is released by the disintegration of a pellet of gunpowder) KZ Zetire used in 37 mm HE A/T Shell (Ref 5, p 558).

38) Mech Imp Fa KZ 36 used in 40 mm HE Shell for Bolero Gun (Ref 5, p 325 & 5, p 563).

39) DA Imp Fa KZ 38; used not indicated (Ref 5, p 561).

40) Mech Imp Fa (self-destructing) KZ 40Z0PWP used in 37 mm HE AA Shell (Ref 5, p 312 & 5, p 557).

41) Graze and DA Fa KZ C/27XLM used in projectiles for Naval Guns (Ref 5, p 565).

42) DA Detonating Type Fa used in 47 mm HE A/T Shell (47 cm Pak Sprg) (Ref 5, p 327 & 5, p 564).

43) Imp Fa (Case Design) M 35ENZ 3/40 used in 47 mm German Ammo (Ref 5, p 331 & 5, p 568).

44) Perc Fa (Skoda Design) used in 75 mm and 83.5 mm projectiles (Ref 5, p 589).

45) Combination Time and Imp Fa VZ 25; used not indicated (Ref 2, p 318).

46) Perc Fa VgZ 36 used in 150 mm Rodded Bomb & 200 mm Sprgr Mortar Bomb (Ref 5, p 589).

47) Mech Imp Fa VgZ 38 used in 50 mm HE Mortar Bomb (Ref 5, p 335 & 5, p 592).

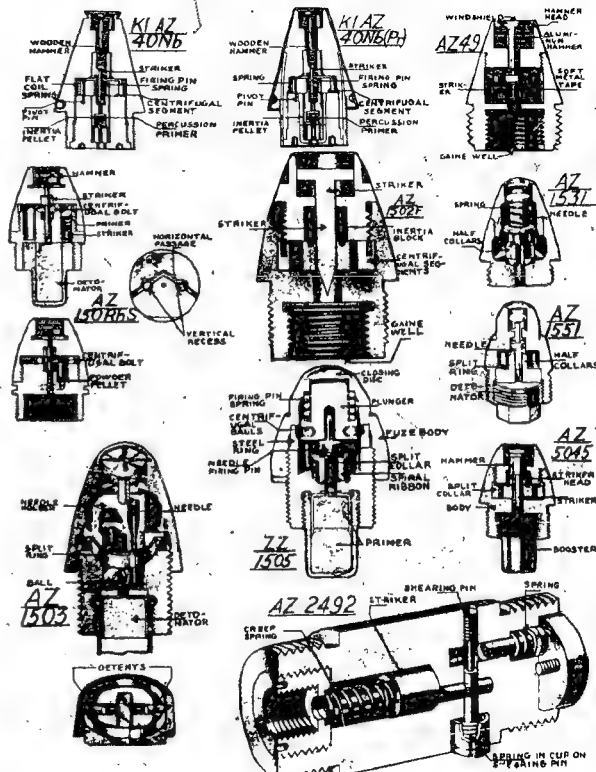
48) Imp Fa VgZ 30 used in 280 mm, 300 mm & 320 mm Rockets (Ref 5, p 397 & 5, p 591).

49) Imp Fa (plastic body) VgZ 3Z ACH used in 80 mm Smoke Mortar Shell (Ref 3, p 381 & 5, p 591).

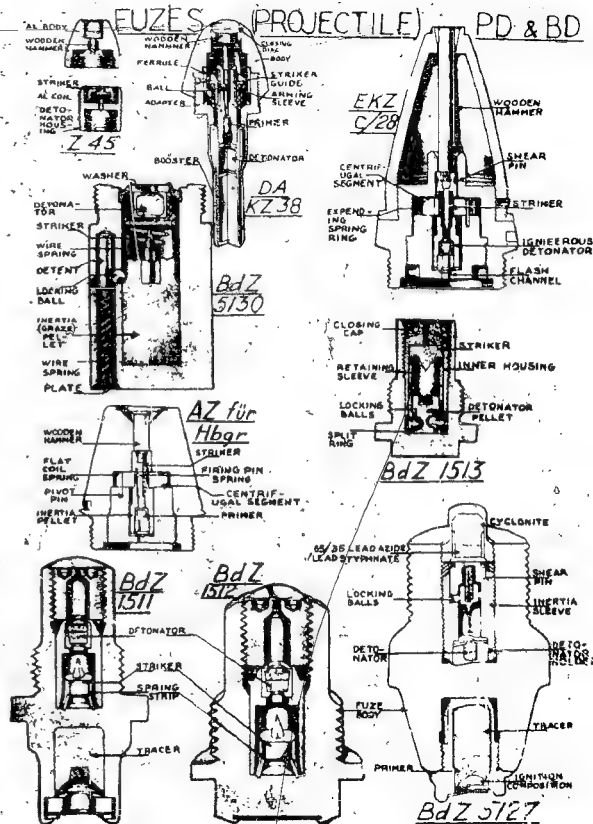
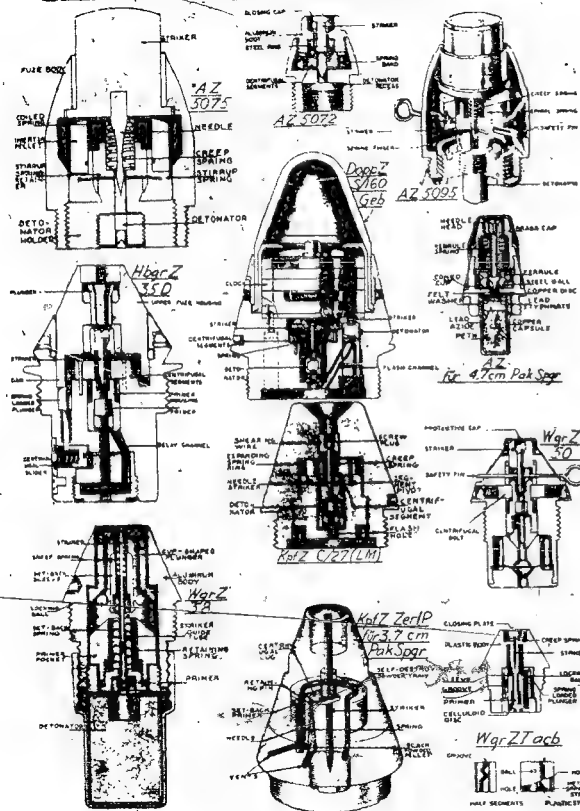
50) Imp Fa Z 45 used in 20 mm Shell (Ref 3, p 304 & 5, p 532).

51) Mech Time Fa Z 25 S/30 & Z 25 S/30gl used in 88 mm & 105 mm HE AA shells (Ref 5, p 555 & 565 and 5, p 554 & 557).

FUZES (PROJECTILE) POINT DETONATING



FUZES (PROJECTILE) POINT DETONATING



Verzögerung" (without delay), while "mv" signify "mit
Verzögerung" (with delay).

- Other German abbreviations are given at the end of this German section following the Vocabulary "American and British Abbreviations"

American and British Abbreviations: AA* Antiaircraft; AC Aircraft; AP Armor-piercing; A/F Antipersonnel; A/T Antitank; B Base; BC Ballistic cap; BDPs Base detonating fuses; C Capped; D Densitizing; DA Direct Action; Blue Electrical; F Fuse; HE High explosive; HG Hollow charge; Imp Impact; Inc Incendiary; M Mark; Mech Mechanical; Perc Percussion.

- Fuze Train (HE Train; Artillery Ammunition Train) (Zündwagen) is described in the general section).

The information in Table 17 is taken from Picatinny Arsenal Technical Report No 1535, pp. 11-15 and some Chem-

"G 3" Enchanted. See Enchanted "G 5".

Guine-Sec Detonators Used in Fuzes.

Galena. See Galose.

Gasdruckpatronen (Gas Pressure Cartridges). See general

section and also the article entitled "Die Entwicklung der Gasdruckpatronen in Deutschland" by E.W. von Horn, in *Zeitschrift für Gasdruckpatronen*, 1934, No. 1, p. 1.

Gaseous Metal Treatment, such as chromizing of iron

or steel articles by the diffusion of chromous chloride vapor at high temperature, is briefly described in BIOS. 31-17-210 (1046) and 3534 (1046).

Gasless Delay Detonators (Electric). German gasless delay

Al or Cu detonator shells (Hüls) having an outside

diameter of 7.20 mm (for Al) and a length ranging from 52.5 to 85 mm (depending on the 'delay' required) were abrasively cleaned and dried before loading.

"Tetryl was loaded first in two increments, a total of 0.7g, to serve as a base charge; this was followed by

an initiating charge of 0.3g of 60/40 L. A. /L. Se. mixture and a perforated (reinforcing) cap all prepared at 25°C.

Note: Tetryl, as well as L. A. and L. St., were previously

After keeping the loaded detonators for 3 days at 30°

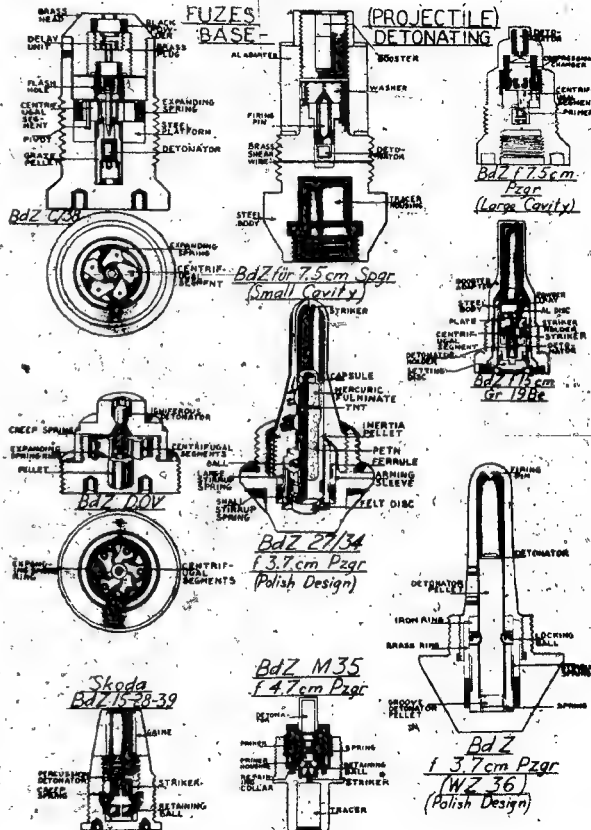
in order to remove all traces of moisture, 50 mg of loose intermediate composition was placed on top

Note: The intermediate composition (powdered mixture of

Sb and KMnO_4) forms a loose connection between the delay composition (to be loaded next) and the initiating composition.

(L A /L St.). The intermediate composition burns with strong flame which facilitates the ignition of L A /L St.

placed in direct contact with L A /L St (Cont'd on p. 65):



Genova Projectile. According to W. Demberger "V-1" Viles (1944), p. 122, Dr Otto Geiseler of Perchleim developed during WW II extremely slender, fiber-stabilized sub-caliber projectiles which could be fired from ordinary gun barrels. It seems that these projectiles were identical with the "arrow projectiles" briefly described under A. These projectiles were used in the 105 mm Antiaircraft Gun (16.5 cm Flak) and in the 280 mm Gun Type 5 (28 cm G.5). It was claimed that by using such projectiles in the Gen-K the range was increased from 37 miles, for the ordinary projectile, to 55 miles with the arrow projectile carrying a sabot behind thick-walled fins. With a lighter type of projectile, which instead of a sabot had an aluminum skirt attached to its middle, a range of about 90 miles was achieved, then using this projectile the tactical dispersion was only about 2 mils. (See also under Arrow Projectile).

GUSTINISPERMETHOFFE (Blasting Explosives)

These are explosive mixtures for blasting purposes. Their composition is not set for purposes cast mines. The following types have been used:

Genstein-Mine. No perchlorate 50, DNN 17, wood meal 3, phosphenite 3 and NG 28 (Ref. 1, p. 123)
Genstein-Dynami. An aluminum 55, TNT 15, K nitrate 5, iron 5 and Na chloride 10; velocity of detonation 4605 m/sec at d 1.17 with a 50 mm diameter confined charge (Ref. 2, p. 192).

Genstein-Kammul. (Genstein-Cammul). A type of commercial explosive aerial rockets of which are given in Table 22.

Table 22 (Genstein-Kammul)

Components and their properties	Designation and source of information			
	Ref. 3, p. 129	Ref. 3, p. 129	Ref. 3, p. 129	Ref. 3, p. 129
No. 1	74.0	85.0	72.0	75.0
No. 2	3.0	8.0	-	-
No. 3	4.0	-	20.0	20.0
No. 4	5.0	5.0	4.0 to 4.0	-
No. 5	2.0	1.0	1.0 to 2.0	1.0 to 2.0
No. 6	8.0	5.0	3.0 to 4.0	3.0 to 4.0
No. 7	-	-	1.0%	1.0%
No. 8	-	-	290cc	280cc
No. 9	-	-	10 cm	20 cm
No. 10	-	-	No 3 cap	No 1 cap
No. 11	-	-	8 cm	8 cm
No. 12	-	-	3000 m/sec	4500 m/sec
No. 13	-	-	1.57	1.46
No. 14	-	-	1219 cal/g	1241 cal/g
No. 15	-	-	3265°C	3300°C

Genstein-Permetol oder Permetol 1. Perchlorate explosive manufactured before WW I by the Sprengstoff A-G. Contains (as in powder and not mine): K perchlorate 50, An nitrate 40, An nitrate 7, TNT 15, flour 4, wood meal 1, and jelly 1%. Its Trauzl test value was 320 Te, pap test 7.0 cm and sensitiveness to impact with a 2 kg weight 70 cm (Ref. 1).
Genstein-Permetol 2 (Genstein-Permetol 2). A type of commercial explosive described in Ref. 3, p. 133. The composition and some properties of these explosives are given in Table 23.

Table 23 (Genstein-Permetol 2)

Components and some properties	Designation	
	No. 1	No. 2
Perchlorate	55	34
Aluminum	43	48
DNN	8	10
DNT	8	10
Carbon (sawdust)	2	2
NG	4	6
Wood meal	4	6
Oxygen Balance, M	-0.3	+1.7
Trauzl Test, cm	530	325

Genstein-Permetol 3 (Genstein-Permetol 3). An minimal type explosive consisting of An nitrate 64.5, DNT 17.0 and Al 3.5% (Ref. 2, p. 114).

- References:
 1) A. M. Shchepkin, Explosives, Chumchik, London, V (1947), p. 364
 2) E. B. Smith, Explosives, Van Nostrand, N.Y. (1939), p. 114
 3) P. H. Smith, Schenck and Sprengstoffe, Stuttgart, Dresden (1937), pp. 129, 133
 4) P. H. Smith, Nitroglycerin etc., Williams & Wilkins, Baltimore (1928), p. 428
 5) C. B. Smith & E. D. Smith, Sprengstoffe und Zündmittel, Springer, Berlin (1936)
 6) T. L. Davis, The Chemistry of Powder and Explosives, Wiley, N.Y. (1943), p. 564.

Genstein-Dynamit (Stratched Dynamite). See under Fill-stoffe.

Genstein (Rifle). See under Weapons.

Genstein 43. German anti-aeromatic rifle, caliber 7.62 mm, developed in 1943. This rifle incorporated some features of a similar Russian weapon, particularly the Degtyarev LMG (light machine gun) and the Tokarev semi-automatic

rifle. The Gewehr 43 weighed 9.75 lbs together with a 0.25 pound sling and a 0.6 pound magazine. [M. Johnson, Jr., Ordnance 29, 306-310, (1943)].

Genstein-Explosivmischungen (Industrial or mining explosives). See Commercial Explosives.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Genstein-Explosivmischungen (Loss of Weight Test) to determine the stability of an explosive or a propellant, is described in Kammul (1944), p. 346 gg.

Glycerol (Glycerin) (Niroglycol, abbreviated to NG). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

Glycerol oder Glycerin (Glycerin, abbreviated to G). See general section.

hazards to a much greater degree than was expected. (See under Reaction of the Base).

The new propellant was called "G" Pulver (G stands for the first fire of Gallwitz).

Due to the fact that "Polyplast" (or straight DEGN) is a good plasticizer for NC, it was possible to prepare propellants more homogeneous than NG propellants and with smoother surface grains. Manufacture of "G" propellants, especially the rolling operation, was much easier and less dangerous and no rolling flaws (often observed in NG propellants) were observed. Another advantage of G propellant was that they guaranteed the incorporation, without becoming brittle, of materials which do not enter part of the plasticization, such as K sulfate (flash reducer, antiragging) (NGs) (See also "Geduldspulver").

Being a good plasticizer, DEGN may be used in smaller quantities than NG and in a wider range. For instance, while content of NG should be 40-45% for optimum results, DEGN may be used in the range of 20 to 45%, the remainder being NC (with such as urethane, or acetone) and one of the following: urethane, phthalates, flash reducers (such as K sulfate or MGs), waxlike, greases, etc.

One such propellant: 61.53% of NC (heat of soluble and insoluble NC giving an average nitrogen content 12.2%), 26.57% of DEGN, 7.55% of chrysotile, 1.60% of urethane, 8.62% of phthalate, 0.25% of K sulfate, 0.1% of graphite and 2.80% of K sulfate had a calorific value of 650-700 kcal/kg as against 820-950 kcal/kg for NG propellants.

As can be mentioned above, the DEGN is more volatile than NG (40% less mass volatile) and is unsuitable for tropical climates.

Inasmuch as the German troops had usually used "G" propellants during the African Campaign, Gen Gallwitz proposed using the simplest product of hydrolysis of (TIG), called "Dyplast" in Germany. This simplest product (TIGDN) was only slightly more volatile than NG (about 1% times) and was quite suitable for hot climates. The replacement of DEGN by TIGDN permitted the production of propellants with even lower calorific value than the ordinary "G" propellants. For instance, one containing 54.55% NC (a heat with an average N content of 12.2%), 25.18 TIGDN, 12.06 urethane, 0.25 MG, 0.50 graphite, and 0.25 K sulfate had a calorific value of 650 kcal/kg. TIGDN possesses the same advantages from the point of view of its plasticizing properties as DEGN and likewise permits the incorporation of plasticizers such as K sulfate and MGs.

"G" propellants are also burning in air efficient in weapons where a projectile remains in the barrel long enough for complete combustion of the propellant. All kinds of gun type weapons and mortars are in this class. All of these weapons have sufficiently long barrels for complete combustion of the powder. "G" propellants in flake form were found unsuitable, however, in medium and small caliber howitzers and mortars because a projectile does not remain for a sufficient time in the barrel for complete combustion of the propellant. In these cases "Geduldspulver" was found to be quite suitable. (See also "Geduldspulver", Reaction of the Base and under "Propellants").

References:

- 1) U. Gallwitz, Die Geschichtsladung (Propelling Charge) Heereswaffenamt, Berlin (1944) (English translation is available).
- 2) G. V. Schickel, et al., General Summary of Explosive Plants,

PB Rep 925 (1945), p 13 and Appendix 9, p 90.
3) H.H. Kipke, Report on Visit to Dinsberg Factory of D A G, CLOS Rep 31-68 (1946), pp 4-5.

GRAMATE (Gr oder g). The term "Gramate" is used in Germany as a base word for various types of rounds. By adding a prefix and/or a suffix to the word the exact nature of the projectile is indicated. E.g.:

Springgranate	Spgr	HE shell
Springgranate 41	Spgr 41	HE shell for improved base gun
Nebelgranate	Nbgr	Smoke shell
Gewehrgranate	Gwgr	Rifle grenade
Handgranate	Hdgr	Hand grenade
Fliegergranate	Flgr	Airborne-dropping (AP) shell
Panzergranate 39	Pgr 39	APCBCHE (Armor piercing capped, ballistic cup, high explosive) shell
Panzergranate 40	Pgr 40	AP shell with a tungsten carbide core
Fliegergranate 41	Flgr 41	AP shell with a tungsten carbide core for tapered bore gun
Gewehrspanzergranate	Gwspgr	Antipersonnel rifle grenade
Gewehrpanzergranate	Gwpggr	Antitank rifle grenade
Gewehrpropagandagranate	Gwpggr	Propaganda rifle grenade
Gewehrballschussler-leucht-granate	Gwblgr	Illuminating parabolic flare projectile
Granate Beten	Gblgr	Antiaircraft shell
Granate Hochladung	Gchlgr	High velocity shell

German Artillery rounds of ammunition may be divided into Panzerammunition and Kammunition:

- 1) Endkammunition, or Endkammunition (One-piece ammunition or round ammunition). It is an one-piece round, the complete round of which may be loaded into the weapon in one operation. This corresponds to American field ammunition. The complete round consists of a cartridge case containing a primer and a propelling charge. The case is permanently crimped to the projectile.

E.g.: Rounds used in AA guns, caliber 30 mm, 28 mm, 30 mm, 37 mm, 40 mm, 42 mm, 50 mm, 75 mm, 88 mm, and 105 mm.

Note: The Germans designated the caliber of guns in centimeters but we designated them in millimeters in order to conform to the American practice.

- 2) Kammunition, or Kammunition (Separated cartridge ammunition) is an ammunition round with intermediate between American round-fired and separate-loading ammunition. It consists of a projectile which is placed into the weapon first and a cartridge case (containing a primer and one or several grains with propelling charge), which is loaded into the barrel afterwards. The cartridge case is not fixed to the projectile. The number of grains with propellant could be varied, according to the range requirement, at the place of firing.

Note: The Germans employed cartridge cases for all their ammunition in order to prevent the escape of gases to the rear of the weapon when the breach is opened; they never

used the rounds corresponding to the American separate loading ammunition.

The Kammunition was used in some 75 mm rounds as well as in 105 mm, 150 mm, 170 mm, 210 mm, 240 mm, 280 mm, and 355 mm guns, or howitzers.

The German Artillery projectiles as well as numerous captured American, Belgian, Czech, Dutch, French, Polish, Rumanian, Russian and Yugoslav projectiles used by the Germans during WW II are briefly described in TM 9-1985-3, pp 519-544. (See under "Captured Projectiles").

Following is the list of these projectiles, arranged by calibers together with the references to TM 9-1985-3.

- 1) 20 mm included: Orlikon AP, Moser AP, Solomou AP, Orlikon HE, Moser HE and Solomou HE are described in TM 9-1985-3, pp 519-544.
- 2) 20/20 mm included: HE 2.0/2.0 mm Spgr/Patr and AP Patr used in Tapered Bore Gun, PzB 41 (pp 371-3).
- 3) 30 mm included: AP, HE, HE-T, AP with Core and Interloaded projectiles used in Solothurn AC Gun (pp 379-42).
- 4) 37 mm included:
 - a) HE-T (3.7 cm Spgr L'apuz) used in Naval C/30 Gun (p 382).
 - b) AP Without Cap (3.7 cm Pgr) used in Pak (p) captured from the Polish (p 382).
 - c) Runded Bomb (3.7 cm Stielgranate, Al) used in Pak 41 (p 383).
 - d) AP Without Cap (3.7 cm Pgr/Patr 18) used in Flak 18 and Flak 36 (p 384).
 - e) HE (3.7 cm Spgr/Patr 40), used in Pak (p 385).
 - f) AP Without Cap (3.7 cm Pgr/Patr) used in Pak (p 386).
 - g) HE (3.7 cm Spgr/Patr) used in Pak (p 386).
 - h) HE (3.7 cm Spgr/Patr C/30) used in C/30 Gun (p 388).
- 40 mm included: HE (4 cm Spgr/Patr) and HE-Lac (4 cm Rr Spgr/Patr) used in Flak 28 (pp 389-9).
- 42/28 mm included:
 - a) HE 4.2/2.8 cm Spgr/Patr L Pak 41 used in L Pak 41 (Tapered Bore Gun) (p 374).
 - b) AP With Core (4.2-2.8 cm Pgr/Patr L Pak 41) used in L Pak 41 (Tapered Bore Gun) (p 374).
- 47 mm included:
 - a) AP With Tungsten Carbide Core Arrowhead Design (4.7 cm Spgr/Patr) used in Czech design improved bore gun Pak (t) and K36 (t) (p 375).
 - b) HE (4.7 cm Spgr/Patr 36) used in some Czech design guns (p 375).
 - c) HE American design (4.7 cm Spgr/Patr (G)) used in Böhler K (3) (p 391).
 - d) APC (4.7 cm Pgr/Patr 36-(t)) used in Czech design gun Flak 37 (t) and Pak (t) (p 392).
- 50 mm included:
 - a) AP With Tungsten Carbide Core Arrowhead Design (5 cm Spgr/Patr 40 KxK) used in the Tank Gun, 5 cm KxK (p 376).
 - b) AP Without Cap (5 cm Pgr/Patr KxK) used in KxK (p 376).
 - c) HE (5 cm Spgr/Patr 38) used in KxK 39 and Pak 38 (p 395).
 - d) APC (5 cm Pgr/Patr KxK) used in the same gun as above (p 395).
 - e) HE-T (5 cm Rr Spgr/Patr 41 L'apuz) used in Flak 41 (p 397).
 - f) HE Mortar projectile used in 3 cm L.G.W. 36" (p 390).

975 mm included:

- a) AP With Tungsten Core Arrowhead Design (7.5 cm Pgr/Patr 41) used in the Antitank Gun, Pak 41 (p 378).
- b) HE (7.5 cm Spgr/Patr KxK 34) and AP With Ballistic Cap and AP Cap (Pgr/Patr 39 KxK 40) used in KxK, KxK 40, SuG 40 and Pak 40 (p 398).
- c) HOC Type 39 (7.5 cm Gr/Patr 39 (H)) used in GebK 15 (p 399).
- d) HOC (7.5 cm Gr/Patr 38 KxK (HL)) used in the KxK, SuG, KxK 40, SuG 40, GebK 36 and the Recoilless Gun for Airborne Troops (LIG 40) (p 400).
- e) HOC (7.5 cm Gr/Patr KxK (HL/B)) used in the same gun as above (p 401).
- f) Smoke (7.5 cm Nbrgr Patr KxK) used in the same gun as above (p 402). (See also Smoke Projectiles).
- g) HE (7.5 cm Gr 40 (15 Aluminum)) used in GebK 15 (p 403).
- h) HOC (7.5 cm lpi) used in LIG 18 and L. Geb IG 18 (p 404).
- i) HOC (7.5 cm lpi 18 AZ/23 ka) used in LIG 18 and L. Geb IG 18 (p 405).
- j) HE (7.5 cm Spgr/Patr 75/50) used in Skoda Dual-Purpose Gun (p 406).
- k) HOC, Type 38 (75 cm Gr/Patr 38 HL/A) used in LFE 18 (p 407).
- l) AP (7.5 cm Pgr/Patr 40 (W) Pak 40) used in Pak 40 (p 408).
- m) APC (7.5 cm Pgr/Patr KxK 38) used in KxK, Spgr, LFK and in Recoilless Gun for Airborne Troops (p 409).
- n) HOC (7.5 cm Gr/Patr 38 HL/A KxK) used in KxK 38, KxK 40, LFK 18, GebK 36, SuG 40, Pak 40, FK 16 and Recoilless Gun 40 (p 409).
- o) APC (7.5 cm Pgr/Patr 39 FES) used in Pak 40, SuG 40 (L/48) and Pak 40; 40/71; 40/72 and 40/74 (p 411).
- p) HOC (7.5 cm Gr/Patr 38 HL/B) used in same gun as given under (a) (p 411).
- q) HE (7.5 cm lpi 38 HL/A) used in LIG 18 and L. GebK 18 (p 423).

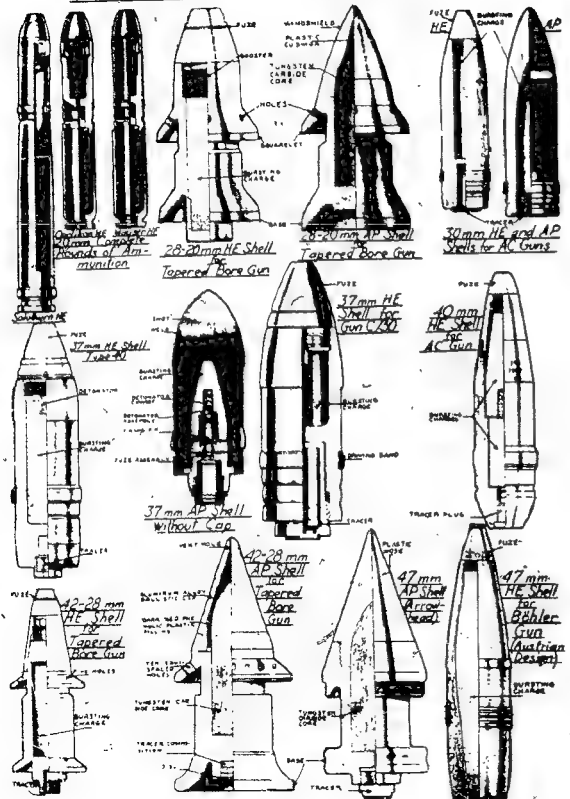
Projectiles used in captured 75 mm Belgian, Dutch, French, Polish and Yugoslav guns not described on pp 410, 413, 415, 419, 420, 421, 423 and 425 of TM 9-1985-3.

10) 75/54 mm was the Brandt Anti-Aircraft projectile developed in France by E. Brandt (p 369).

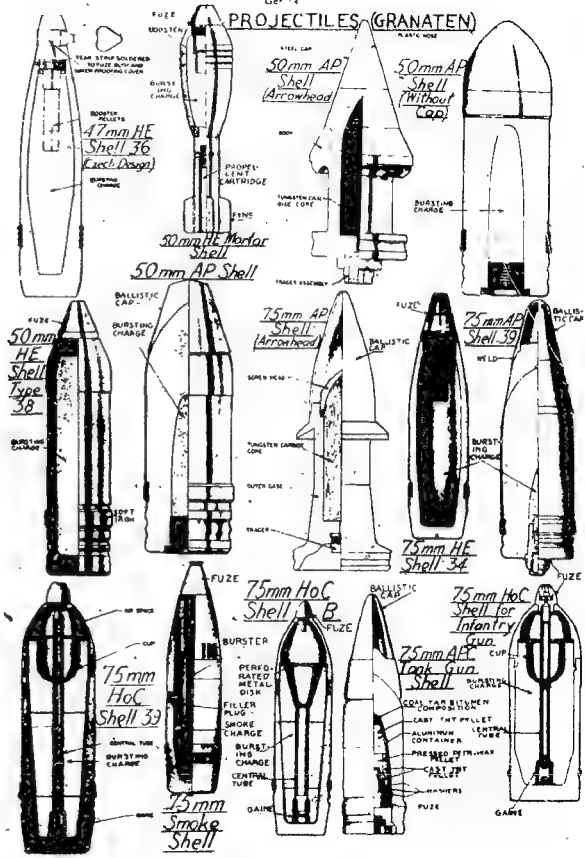
11) 75.8 mm included the following projectiles used in captured Russian weapons:

- a) HE (7.62 cm Spgr 284/4) used in GebK 307 (t) (p 426).
- b) HE (7.62 cm Spgr/Patr 39) used in FK 36 (t) (p 426).
- c) AP (7.62 cm Pgr/Patr 40) used in FK 296 (t) FK 36 (t) and Pak 36 (t) (p 427).
- d) APC (7.62 cm Pgr/Patr 39 rot) used in Pak 36 (t) (p 428).
- e) HE (7.62 cm Spgr 280/2) used in JKH 290 (t) (p 429).
- f) HE (7.62 cm Spgr 284/4) used in GebK 307 (t) (p 430).
- g) HOC (7.62 cm Gr 38/2 HL/B) used in JKH 290 (t) (p 430).
- h) HE (7.62 cm Spgr 39/2) used in JKH 290 (t) (p 430).

PROJECTILES (GRANATEN)



PROJECTILES (GRANATEN)



12) 76.5 mm projectiles were used in captured Austrians, Czechs and Yugoslavs 7.65 cm weapons (pp 432-433)

13) 80 mm included:

- a) HE Mortar proj (8 cm Wgr 36 and Wgr 39) used in WGr 34 (p 529)
- b) Colored Smoke proj (8 cm Wgr 36 Dwe) used in WGr 34 (p 535). (See also Smoke Projectiles)
- c) HE Smoke proj (8 cm Wgr 34 Nb) used in Mortar, MGW 34 and KxWg 42 (p 532)
- d) 82.5 mm included: 8.5 cm Sprg (r) and Gr 23/20 (r) used in captured Czech AA Gun, Flak M/2 (r) (pp 456-7)

15) 88 mm included:

- a) APC (8.8 cm Sprg Patr 39) used in Flak 41 (p 438)
- b) HE (8.8 cm Sprg Patr L/4.5 (Ka)) used in Flak 18, Flak 36 and Flak 37 (p 438)
- c) AP (8.8 cm Sprg 41) used in Flak 36 and Flak 41 (p 439)
- d) AP with Tungsten Carbide Core, Type 40 (8.8 cm Sprg 40) used in Flak 36 and Flak 41 (p 439)
- e) HE (8.8 cm Sprg Patr L/4.7 FES) used in Flak 41 and Flak 43 (p 441)
- f) APC (8.8 cm Sprg Patr mB42) used in Flak 18, Flak 36 and Flak 37 (p 441)
- g) HE, Type 43 (8.8 cm Sprg Patr 43) used in KxK 43, Stuk 43 (L/71) and Pak 43 and 43/71 (L/71) (p 442)
- h) HE (8.8 cm Sprg 39/43) used in Pak 43 and Pak 43/41 (p 442)
- i) HE (8.8 cm Sprg Flak 41) used in Flak 41 (p 443)
- j) HoC (8.8 cm Sprg HL) used in KxK 36 (L/56) (p 444)
- k) HE (8.8 cm Sprg L/4.5) used in KxK 36, Flak 18, Flak 36, Flak 37 and in Modified Russian AA Gun W/3.8 cm Flak 39 (r) (p 444)

- l) HE, with Controlled Fragmentation (8.8 cm Sprg L/4.5 ZGZ) used in KxK 36 (L/56) (p 445)
- m) AP (8.8 cm Sprg) used in Flak 18, 36, 37 and in Flak 39 (r) (p 446)
- n) AP (8.8 cm Sprg 39) used in Pak 43, Pak 43/41 (L/71) and Stuk 43 (L/71) Self-propelled gun (p 446)

- o) AP (8.8 cm Sprg 39) used in Flak 18, 36 & 37, KxK 36 (L/56) and in Flak 39 (r) (p 448)
- p) Incendiary Shrapnel (8.8 cm Gr Br Schr Flak) used in Flak 18, 36 and 37 (p 448)

16) 90 mm included:

- a) HoC proj Type HL/B and Type HL/C are described in TM 9-1985-3, pp 450-1, but their uses are not given
- b) HE Czech proj [10 cm DoppZGr M 2110] used in captured Czech, Polish and Yugoslav Light Field Howitzer (p 451)

- c) HE Yugoslav proj [10 cm Sprg DoppZ 311 (r) and Sprg (A2) 310 (r)] used in captured Czech, Polish & Yugoslav Light Field Howitzers and Model 28 Yugoslav Mountain Howitzer (p 452)

- d) HE Czech proj [10 cm DoppZGr 30 (r)] used in Czech, Polish and Yugoslav Light Field Howitzers (p 453)

- e) HE Polish proj [10 cm Sprg (r)] used in Czech, Polish and Yugoslav Light Field Howitzers (p 455)
- f) HE German proj (10 cm Sprg 38) used in Czech, Polish and Yugoslav Light Field Howitzers (p 454)
- g) HE Mortar proj (10 cm Wgr 37) used in NWG 35 (p 535)

17) 90.5 mm included:

- a) HE (10 cm Gr 19) used in K 18 (p 456)
- b) HE used in K 17, O1 nA and K 17 (p 457)

c) AP used in several Light Field Howitzers (pp 457 and 459)

- d) HE (10 cm Sprg L/4.4) used in Flak 38 (p 467)
- e) AP-T (10 cm Sprg rtr) used in Flak 38, Flak 39, KxK 18 and KxK 16

- f) AP (10 cm Sprg rtr) used in Light Field Howitzer (LFH 16) (p 470)

- g) HE used in Light Field Howitzer LFH 16 (p 471)

- h) Smoke used in Howitzers (LFH 16, LFH 18, LFH 18MB and StuH 42) (p 472)

- i) HE for Long Distance Use in Light Field Howitzers 18 with Axial Brake (LFH 18MB) (p 473)

- j) HoC Type A, HoC Type B and HoC Type C used in the same Light Field Howitzers as listed under (h) (pp 474-77)

- k) HE, Model 15, Model 73 and Model 28 used in the 10 cm Shoda Howitzer (p 477-60)

- l) HE (10 cm Sprg Patr L/4.4 Ka) used in Flak 38 and Flak 39 (p 480)

- m) HE (10 cm Sprg 19 Ka 15) used in K 18, KxK 18 and KxK 17 (p 481)

- n) HE proj with disintegrating band is described briefly on p 369 of TM 9-1985-3

- o) Projectiles used in captured 105 mm Belgian, French, Polish, Russian and Yugoslav weapons described on pp 459, 461 and 463-467 of TM 9-1985-3

- p) HE (10 cm FHGrGr mR 11) used in Light Field Howitzers: 18, FH 18/1, FH 18/2, FH 18 mm, FH 18/33 and FH 18/49 (p 536)

- q) 122 mm included HE projectile 12.2 cm Sprg FEW (r) used in captured Russian guns K 390/1 (r) and K 390/2 (r) (p 481)

- r) 128 mm included:

- a) HE (12.8 cm Sprg Patr L/4.5), described briefly on p 482

- b) AP (12.8 cm Sprg FES) used in Flak 40 (p 483)

- c) AP (12.8 cm KPS) used in Flak 40 (p 483)

- d) AP (12.8 cm Sprg 43) used in Flak 44, self-propelled (p 484)

- 20) 150 mm included:

- a) HE With Disintegrating Bands, Sabot Type (p 570)

- b) HE (15 cm AZGr 37 (r)) used in Czech Medium Howitzer sFH 23 (r) (p 485)

- c) HE (15 cm CGr 42) used in K 18 (p 486)

- d) HoC (15 cm Jgr 39 HL/A) used in StuH 4XL/12) and sJG 33 (p 486)

- e) A/C (15 cm Gr 19 rot Be) used in K 18 and K 39 (p 487)

- f) Czech projectiles, such as 15 cm GmM 25 (r) (p 488), 15 cm AZGm 34 (r) (p 488), 15 cm MinGr M 13/19 (r) (p 489), 15 cm MinGr 28 (r) and 15 cm MinGr M 28 (r) (p 490) used in captured Czech Field Howitzers

- g) HE (15 cm Jgr 38 FES) used in the Assault Howitzer StuH 43 (p 491)

- h) AP (15 cm PaSprg L/37 mHBe) used in K 18 (p 491)

- i) HE (15 cm Gr 36 FES) used in sFH 18 (p 492)

- j) HoC (15 cm Gr 19 HL) used in sFH 18 and sFH 13 (p 492)

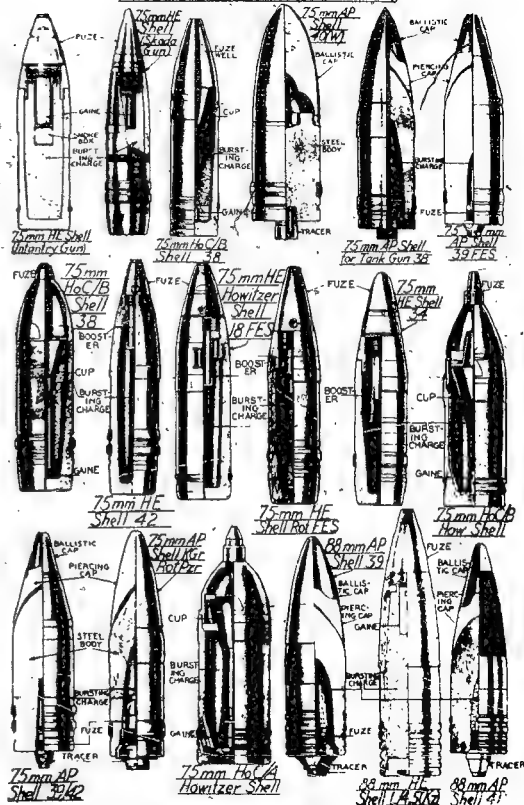
- k) A/C (15 cm Gr 19 rot Be) used in K 18, K 39 and in K (E) (p 493)

- l) HE (15 cm Gr 19mZdg 36) used in sFH 18 (p 494)

- m) HE proj of cast steel (15 cm Gr 19 Stg) used in sFH 18, sFH 15 and sHT (p 495)

- n) Smoke (15 cm Gr 19 NB) used in sFH and sFH 13 (p 497)

PROJECTILES (GRANATEN)



Ger 77



Ger 78



p) Rodded Bomb (15 cm Stielgranate 47) used
aIG 33 (p 498)

c) HE (15 cm Gr 18) used in aFH 13 (p 500); HE (15 cm Jgr 38) used in aIG 33; HE with Base Fuse and Ballistic Cap (15 cm Spgr L/4.4 BaZ mix Haube) used in Ki Men Lal (p 504); HE with None Fuse (15 cm Spgr L/4.6 Kz) used in K39 (p 504).

- u) SAP (15 cm Hpagr) used in K 39 (p 504)
- t) AP (15 cm Pgr) used in K 39 (p 504)
- v) Smoke (15 cm Gr 38 Nb) used in uFH 18 (p 506)
- v) A/C (15 cm Gr 19 Be) used in uFH 18 (p 507)

21) 152 mm included the following types used in
the Vietnam theater:

a) HE (15.2 cm Sgr 436) used in KH 433/1 (r)
and KH 433/2 (r) (p 510)
b) A/C (15.2 cm Gr 434 Be) used in the same
weapons as above (p 511)

a) HE [15.5 cm StGr 422 (f)] used in K 418 (f)
K 419 (f) and K 420 (f) (p 512)

c) HE [15.5 cm Gr 417 (I) and 1mmgs 415 (I) used in sFH 414 (I) and sFH 17 (p; p. 513-4)
d) HE [15.5 cm Gr 421 (I)] used in 15.5 cm K 420 (I) (p 515)

23) 170 mm included:
a) HE (17 cm KGr 38Hb) used in Ki Mrs Laf (p 316)
b) HE (17 cm KGr 39) used in Ki Mrs Laf (p 517)
24) 194 mm included the HE proj [19.4 cm StG G
486 (21) used in captured French Railroad Gun

25) 200 mm included the NE Mortar Projectile
20 cm Wgr 40 (p 514)

b) Flare projectile (20.3 cm Leuchtgrt) used i
K(E) (p 520) (See under Flares) .

c) HE (20.3 cm Spgr L/14 Kz (Hb) and Spgr L/4.7 Kz mHb used in K(E) (p 521)
d) SAP (20.3 cm Spgr L/4.7 BdZ mHb)' used in K(E) (p 520)
27) 210 mm included A/C proj (21 cm Gr 19 Be) w

28) 240 mm included:
a) HE (24 cm Spgr 1/4.5 BdZ mHb and Spgr 1/4.2 mHb) used in Theodor Bruno Railways Gun, TbBrK(E) (p 524)

29) 280 mm included:
a) Rifled 28 cm projectile. Its nomenclature and uses are unknown (p 326)

30) 355 mm (p 3294) (its caliber was also given as 355 mm) howitzer M1934 (its caliber was also given as 355 mm) 355 mm included HE mortar proj (38 cm GrBe) and HE Rocket Assisted Rifled proj (28 cm R 413 and Gr 353 used in K 5 (E) (p 327-28)

and Smoke, 10.133 cm wgr 40 Nb/p 33%

Armor-piercing, capped; HE High-explosive; HoC Hollow charge; Inc Incendary; SAP Semi-armor-piercing; T Tracer
German Abbreviations: See Abbreviations at the end of this German section.³
References: Anon, Technical Manual TM 9-1985-3 (1953), pp

The same information is given in the following references:

- 1) Anon. Enemy War Materials Inventory List, Ammunition, Supreme Headquarters AEF, (1945), pp 1-154
- 2) Anon. Recognition Handbook of German Ammunition, 1943

Supreme Headquarters AEF (1945)
3) Amos, German Artillery Projectiles and Fuzes, Ordnance
Navy Bomb Disposal Center Aberdeen Proving Ground and U.S.
Navy Bomb Disposal School, pp 1-177 (No date).
Note: According to Ref 1, pp 131-3, the following larger

caliber projectiles were used by the Germans: 380 mm HE and AP for 38 cm Siegfried Kanone C/34; 406 mm HE and AP for 40.6 cm Adolf Kanone or for Navy gun, Schiffskanone C/34; 420 mm HE, Anticoacete for 42 cm howitzer, called *Karl* *Mörser*; 540 mm HE for 54 cm heavy howitzer, called *Karl Mörser*; 615 mm HE for 61.5 cm

The following types of grenades are described in

1) Stick Hand Grenades, Modela 24, 39 und 43 (Stiehlhandgranaten 24, 39 und 43) (pp 319-20)

3) Shaving Stick Offensive Hand Grenade (p 322)
4) Magnetic Antitank Hand Grenade, 3kg. (Hafthohlladungsgranate, 3kg) (p 323) (See *Hafthohlladung*)
5) Hollow Charge Stick Type Hand Grenade (p 324)

6) Antitank (Hollow Charge) Hand Grenade (Panzerwurfmine) (p 324)
7) Smoke Hand Grenades, Models 39 and 41 (Nebelhandgranaten 39 und 41) (pp 325-6)
8) Smoke Hand Grenade 14 (Blaskörper 14) (p 327)

9) Smoke Hand Grenade 24 (Blendkörper 24) (p 328)
10) Smoke Hand Grenade, Egg Type (p 329)
11) Hand Smoke Signal, Red (Hundrauchzeichen-
Rot) (p 329)
12) Lacrimatory Hand Grenade (Tear Bomb) (p 330)

14) 61 mm Antitank (Hollow Charge) Rifle Grenade (S.S.Gewehrpanzergranate, 61 mm) (Two types, pp 331 and 332)

16)* Antitank (Hollow charge) Rifle Grenade (Gewehr Panzergranate) (p 334)

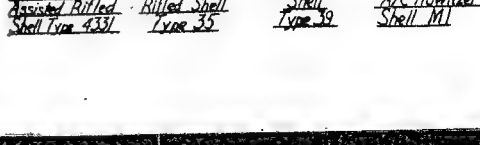
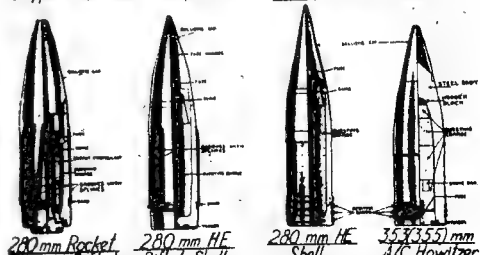
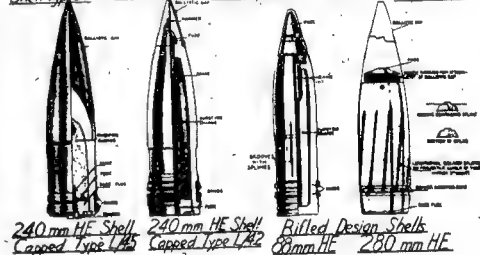
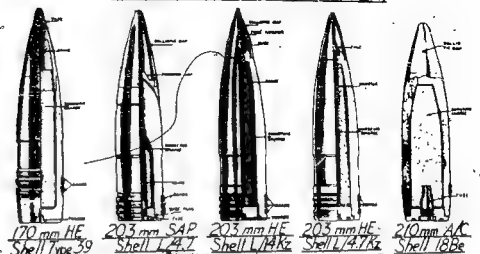
18) Large Antitank (Hollow Charge) Rifle Grenade
(Grosse Gewehr Panzergranate) (p 336)

20) Propaganda Rifle Grenade (Gewehr Propagandagranaat) (p 338)
21) Illuminating Parachute Rifle Grenade (Gewehr Fallschirmleuchtgranaat) (p 339)

22) Hollow Charge Grenade, called Faustpatron (p 339)
23) Pistol Grenade (Wurfkörper Leuchtpistole) (p 340)
24) 27 mm Pistol Grenade HE Egg Type, fired from

24727 mm Pistol: German H&K L88 type, third model
x Walther pistol (p 341)

PROJECTILES (GRANATEN)



Other successful guided missiles were:

- a) Schmetterling (Butterfly), also known as the Hs-117 (Ref 2, p 35)

Note: He is an abbreviation for Henschel, the name of the builder.

- c) Rheintochter (Daughter of the Rhine) series such as Rheintochter I, II and III (Ref 2, p 40)
- d) Enzian (Gentian, a species of blue flower) series, ranging from E-1 to E-5 (Ref 2, p 43 Ref 3, p 59)

e) Feuerlilie (Fire Lily) series, of which the Hecht (pike) was one of the first successful. T-Stoff and Z-Stoff were used in it. The Hecht was succeeded by the Feuerlilie F-25. The last of the series was the H-25, used only for research (Ref 2, pp 43-47, Ref 3, pp 29-40)

- f) Bachem HFP-20 Master (Viper) (Ref 2, p 47)
- g) Ruhrstahl (Steel-Birch) the Ruhr series ran from X-1 to X-7, of which the X-4 was the most important (Ref 2, p 36 and Ref 3, pp 30-2)

h) Ma (Hüschel), the name of builder) series, including the previously mentioned Hs-117 (Schmetterling), as well as Hs-117H, Hs-293, Hs-294, Hs-295, Hs-296 and Hs-298 (Ref 2, pp 52-54 & 54-60, Ref 3, pp 32-33)

- i) Fritz X (FX-1409), a glide bomb (Ref 2, p 55)
- j) Bachem Apache - an odd-looking guided missile (Ref 2, pp 61-62)
- k) BV-246 (Ref 2, p 63)

l) V-2, is briefly described separately under V-2.

It could be launched as a guided missile.

- m) Antipodal Bomber (Ref 4, pp 57-58)

n) Tu-16, a liquid rocket (Ref 5, p 223).

References:

- 1) Anon, Army Ordnance 31, pp 28-30 & 121-24 (1946)
- 2) F. Ross, Jr., Guided Missiles, Rockets and Torpedoes, Lothrop, Lee & Shepard Co. Inc, N.Y. (1951), pp 14-66

3) A. Dacrocq, Les Armes Secrètes Allemandes, Berger-Levrault, Paris (1947) pp 90-99

4) K.W. Gailard, Development of the Guided Missile, "Flight" Publication, London (1952), pp 2-19, 47 & 49-59

5) Anon, Dept of the Army Technical Manual TM 9-1985-2 (1953), pp 195-233

Note: Additional information on guided missiles, also called Directed Missiles, may be found in the following CIO Reports: 28-Mc, 29-41, 31-13 and 32-66, which were published in 1945 and 1946

(See also Great Enzian Guided Missile, Rockets and V-2)

Gummidynamit. A rubberlike elastic explosive mass obtained on dissolving collodion cotton in NG. This is called also Sprenggelatine (Blasting gelatin).

Gvn (Geschütz). See Cannon and also Weapons.

Guncotton-Dynamit. See Triaxial Dynamit.

H. One of the abbreviations for Hexogen or Hexo (Cyclonite).

H₈, H₁₀ etc. Hexogen phlegmatized with 8%, 10% etc Montan wax.

H-1, H-2, H-5, H-8 Explosives, German Ammonites,

described under Ersatzsprengstoffe.

HA. One of the abbreviations for mixture of RDX (Hexogen) and Al (aluminum).

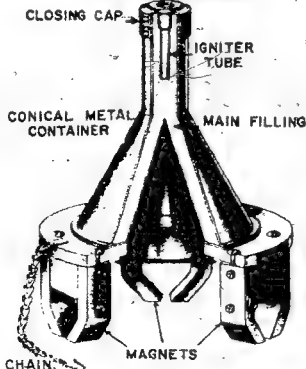
Haft-Hohlladung (Adhering or Sticking Hollow Charge). One of the devices consisted of a conical metallic container (filled with 5 lb 5 oz of a HE) to which was attached an elongated Apex, serving as a hand grip and containing the igniter pellet (PETN/Wax) and a pull (friction) delay igniter (4; or 7 seconds). Attached to the base of the conical section was a plywood frame-work carrying three powerful horseshoe magnets. A heavy chain with a hook was attached to the framework. Total weight 3 kg.

The device could be used either as a hand grenade or as a land mine. In the first case the cord of the friction igniter was pulled off and the grenade thrown against the approaching vehicle. In the second case, the device was buried in the ground, close to the surface, with the magnets up and with the igniter cord attached to the ground. As the approach of a vehicle the magnetic attraction caused the grenade to jump towards some iron or steel part and attach itself to it. Simultaneously the cord was pulled, thus setting off the explosive, main consisting of delay igniter, explosive and main charge. (Ref 2, 11) It was claimed that this charge could penetrate as much as 110 mm of armor. (Ref 1, pp 32-4).

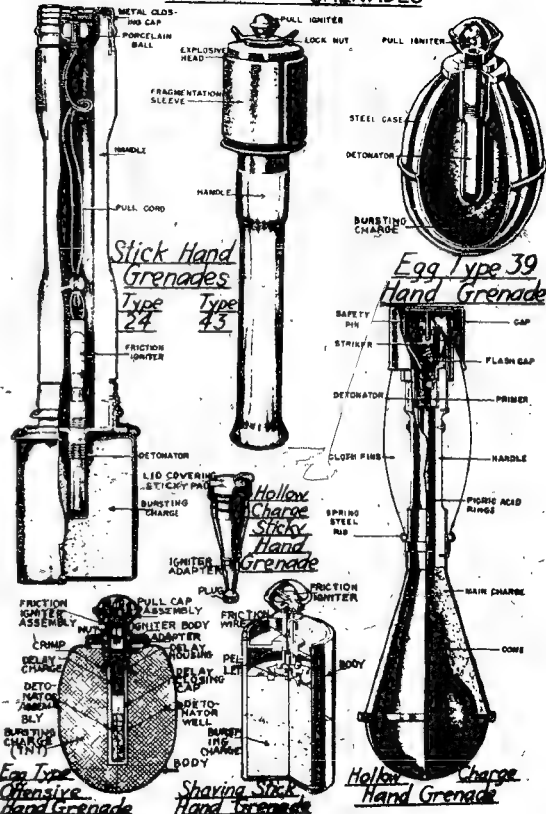
Another magnetic antitank charge is described in Ref 1, pp 30-3 under the name of Panzerhund 3. It consisted of a bottle-shaped cardboard container with 2 1/3 lb of hollow charge (TNT or RDX/TNT). Three pairs of magnets were mounted at the bottom of the bottle, and a 7" sec friction igniter was located in the neck of the bottle. Total weight of the device was 8 lb.

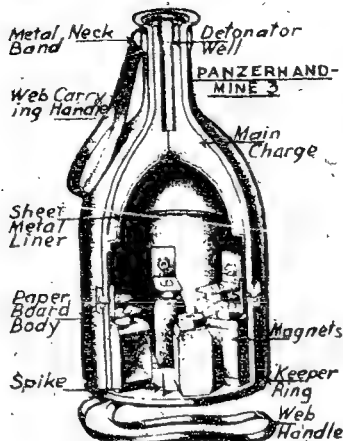
The device was apparently designed to be placed by hand on the tank and the igniter pulled after it has been positioned. If the target was of non-magnetic material such as wood, the charge could be attached by means of 3 spikes located at the bottom of the device. (pp 262-3).

HAFTHÖHLADUNG



HAND GRENADES





In another type of adhering (sticking) antitank hollow charge there were no magnets but a sticky pad (located at the wide part of the conical body) served for attaching the charge to a tank (Ref 1, p 374).

- References:
 1) Dept of the Army Tech Manual TM 9-1985-2 (1953), PP 302-3 & 313-4
 2) H.H. Bullock, Picatinny Arsenal; private communication.

Thermite (Adhering Mine). An antitank hollow charge device consisting of a conical container (filled with HE), provided with a Cat mop and a handle. The wide portion of the cone was covered with a layer of a low melting colloidal plastic resin (p. ca 500) retained on the surface by means of an open mesh cloth. In back of the flat top, which consisted of sheet metal, was placed a thermite-type charge ($Mg + Al + KClO_3$) and in back of the latter a time fuse. The operator hid in a hole and, at the approach of the tank, ignited the fuse which, in turn, ignited the thermite. Just as soon as the heat of the thermite melted the resin, the device was struck (by the operator) to the bottom armor plate of the tank. At the same time the heat of the thermite set off the detonator and this in turn initiated the main charge.

This device was in an experimental stage when the war terminated.
 Reference: E.E. Richardson et al, CIOS Rept 25-18 (1945), pp 25-3.

Makhlissit. Same as Perokhlantiv.

Makhlissit oder Lagerbestelligkeit (Stability in Storage) See in the general section.

Handfeuerwaffen (Small Arms) See under Weapons.

Handabzugsscharenprengstoffe (Explosives Safe to Handle and to Transport) See Davis (1943), p 347.

Hornstoff (Urea). See general section.

HC Mixture. A smoke mixture consisting of hexachloroethane and powdered zinc.
 Reference: Anon, Field Artillery Journal B, 352-3 (1943).

Heavy A/T Mine. See under Landmines and also on pp 263-7 of TM 9-1985-2 (1953).

Hahelkinder (Lever Type or Schuko Igniter). See Pressure Igniter under Igniter.

Hoch Guided Mine. See Pike (Tech) Mine.

Hellhoff Explosives According to Ger P 12,122 of 1880, it was prepared by the nitration of purified turpentine, followed by washing, drying and mixing of the nitroar with oxygen carriers, such as K (or Na) nitrate (or chlorate), etc. It was claimed that this explosive mixture was very powerful.
 Reference: See under Hellhoffit.

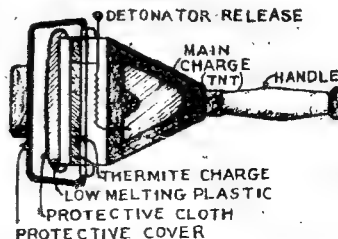
Hellhoffit (Hellhoffite). One of the Sprengel type explosives, invented about 1870 by Hellhoff and Grünig. It consisted of 28 parts of nitrobenzene and 72 parts of fuming nitric acid. This liquid was sometimes used absorbed on kieselgur (see Gubzhellhoffit). The disadvantage of these Sprengel type explosives was their extreme corrosiveness (Ref 1).

According to Thorpe (Ref 2), Hellhoffit was tried in shells, the two ingredients being mixed during flight exploded on impact (see also Anilite under French explosives).

Stettbacher (Ref 3 and 4) investigated Hellhoffit and its modifications and found that the glass-lined depth charges (Tiefenbomben) containing Hellhoffit, were much more effective than those loaded with picric acid. The mixture consisting of fuming nitric acid (d 1.52) 64.51, nitrobenzene 25.81 carbon disulfide 6.45 and aluminum bronze 3.25% was found to be one of the most effective. A mixture prepared by dissolving 66.7 parts of dinitrobenzene in 100 parts of fuming nitric acid was also claimed to be effective.
 Reference:

- 1) Davis (1943), p 354 2) Thorpe's Dictionary, v 4 (1940), p 545 3) A. Stettbacher, S S 38, 158 (1943)
- 4) A. Stettbacher, Spreng- u. Schiessstoffe, Zürich (1948), p 71.

HAFTMINE



Hemgels. Smokeless propellant was based on nitrocellulose pulped with some chemicals as dextrinate (1907), p 273.

Henschell or He. A guided missile, see under Missiles.

Herkulin of Dyckerhoff. An explosive made in a concentrated aqueous solution of picric acid and Am product was dried and mixed with pulverized sulfur and K, or Na nitrate. Reference: L.Gody, Traité des Explosifs (1907), p 551.

Hetzler (Baiter). A Czech design Destroyer, Jagdpanzer 38 (t) (See Destroyer).

Hausgeschosse (Grasshopper). A (Waffenkörper) such as for 105 mm German early in the WW I. The of the Illustrated Record of Germany, 1945, War Office, London (1947). Note: The above British books fear that they are 'confidential' with British sources.

Hexo, Hexamin, Hexamitridiphenyl (Hexanitrodiphenylamine) (HNDPA) general section under Diphenyl information concerning the Hexa in Germany during WW II.

At Allendörfer Fabrik of manufacture was as follows:
 To a charge of 1000 kg of in a V2A stainless steel capacity (fitted with an agitator, a cooling jacket and a diphenylamine was added temperature was maintained with weak of 30-40°. The precipitated oil, washed thoroughly spread and packed.

HNDPA was used by the of R 1 in an underwater explosion 40 and TNT 85%. During WW replaced by the one containing 53.7 and Al 16.4%. Another contained HNDPA 23.0, T Stettbacher (Ref 5) cites a HNDPA with 30-98 TNT and 16% Schiesswolle 18, TSNV-1-101 Reference:

- 1) A. Stettbacher, Procar (Switzerland)
- 2) U.S. Naval, Tech Mission 515-45, Hexanitrodiphenylamine, PB Rept 38, 154 (1945)
- 3) O.W. Strickland et al, PB Re 4) Anon, Allied and Enemy Proving Ground, Md (1946)
- 5) A. Stettbacher, Spreng- (1948), pp 78-79.

Hexa 5-22, 5-24 and E-4. Ger containing hexanitrodiphenyl: Ersatzsprengstoffe.

Hexastoff (Urean). See general section.

HC Mixture. A smoke mixture consisting of hexachloroethane and powdered zinc.

Reference: Anon, Field Artillery Journal B, 352-3 (1943).

Heavy A/T Mine. See under Landminen and also on pp 265-7 of TM 9-1985-2 (1953).

Helsänder (Lever Type or Schuko Igniter). See Pressure Igniter under Igniter.

Hecht Guided Missile. See Pike (Hecht) Missile.

Heilhoff Explosive. According to Ger P 12,122 of 1880, it was prepared by the nitration of purified tar oil, followed by washing, drying and mixing of the nitrotar with oxygen carriers, such as K (or Na) nitrate (or chlorate), etc. It was claimed that this explosive mixture was very powerful.

Reference: See under Heilhoff.

Heilhoff (Heilhoffite). One of the Sprengel type explosives, invented about 1870 by Heilhoff and Grünke. It consisted of 28 parts of nitrobenzene and 72 parts of fuming nitric acid. This liquid was sometimes used absorbed on kieselguhr (see Gubkheilhoff). The disadvantage of these Sprengel type explosives was their extreme corrosiveness (Ref 1).

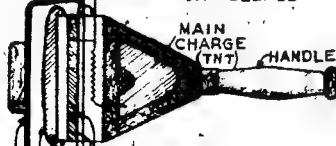
According to Thorpe (Ref 2), Heilhoff was tried in shells, the two ingredients being mixed during flight exploded on impact (see also Anilind under French explosives).

Sternbacher (Refs 3 and 4) investigated Heilhoffite and its modifications and found that the glass-lined depth charges (Tiefenbomben) containing Heilhoffite, were much more effective than those loaded with picric acid. The mixture consisting of fuming nitric acid (4 1/2%) 64.5%, nitrobenzene 25.81%, carbon disulfide 6.45% and aluminum bronze 3.23% was found to be one of the most effective. A mixture prep by dissolving 66.7 parts of dinitrobenzene in 100 parts of fuming nitric acid was also claimed to be effective.

References:
1) Davis (1943), p 354
2) Thorpe's Dictionary, v 4 (1940), p 543
3) A. Sternbacher, S 5 38, 158 (1943)
4) A. Sternbacher, Spreng- und Schießstoffe, Zürich (1948), p 71.

HEXAMINE

DETONATOR RELEASE



THERMITING CHARGE
LOW MELTING PLASTIC
PROTECTIVE CLOTH
PROTECTIVE COVER

Hexanit. Smokeless propellant, patented in 1885; was based on nitrocellulose straw previously treated with some chemicals as described in Daniel, Dictionnaire (1902), p 733.

Hexschell or Hs. A guided missile (q v) developed during WW II.

Hexolin of Dyckerhoff. An explosive prep by soaking sawdust in a concentrate aqueous solution of equal parts of picric acid and Am nitrate. The resulting product was dried and mixed with various amounts of pulverized sulfur and K, or Na nitrate.

Reference: R. Goulet, Traité des Matières Explosives, Nanterre (1907), p 554.

Hetzer (Baier). A Czech designed and constructed Tank Destroyer, Jagdpanzer 38 (t) (See under Panzer).

Hexachoke (Graessbopper). A series of weapon carriers (Kaffenzüger) such as the 105 mm Gun, developed by the Germans early in the WW II. They are described in vol III of the Illustrated Record of German Army Equipment 1919-1945, War Office, London (1947).

Note: The above British names were not consulted for fear that they are "confidential" or "secret" as is usual with British sources.

Hexa, Hexamina, Hexamitridiphenylamin, oder Hexyl (Hexamitridiphenylamin) (HNDPPha). Described in the general section under Diphenylamine. The following information concerning the manufacture and use of Hexa in Germany during WW II is available:

At Allendorf Fabrik of VASA-G, the method of manufacture was as follows:

To a charge of 1000 kg of 99% nitric acid placed in a V2A stainless steel mixer of 2 cubic meter capacity (fitted with agitator rotating at 60 RPM, a cooling jacket and cooling coils) 300 kg of diphenylamine was added gradually while the temperature was maintained at 90°. The solution was filtered with weak nitric acid and cooled to 30-40°. The precipitated HNDPPha was filtered off, washed thoroughly with water, then dried, screened and packed.

HNDPPha was used by the Germans at the start of WW II in an underwater explosive containing HNDPPha 40% and TNT 50%. During WW II, this explosive was replaced by the one containing HNDPPha 27.9%, TNT 55.7% and Al 16.4%. Another underwater explosive contained HNDPPha 25.0%, TNT 61.8% and Al 15.2%. Sternbacher (Ref 3) cites a mixture consisting of HNDPPha with 30-40% TNT and 16% Al (See also Hexamine, Schießstoffe 18, TSMV-1-101 and Ersatzsprengstoffe). References:

- 1) A. Sternbacher, Prosser (Switzerland), 33-45 (1943)
- 2) U.S. Naval Tech Mission in Europe, Tech Rept 513-45, Hexamitridiphenylamine Manufacture in Germany, PB Rept 38, 154 (1945)
- 3) O.W. Stickland et al, PB Rept 1820 (1943), pp 13-17
- 4) Anon, Allied and Enemy Explosives, Aberdeen Proving Ground, Md (1946)
- 5) A. Sternbacher, Spreng- und Schießstoffe, Zürich (1948), pp 78-79.

Hexa 5-22, 5-26 and E-4. German substitute explosives containing hexamitridiphenylamine described under Ersatzsprengstoffe.

Hexodi—German name for Hexamethylenetetramine Dinitrate, $C_6H_{12}N_4 \cdot 2H_2O$ (See KA-Verfahren under Hexogen).

Hexol. An explosive mixture consisting of 75% Hexogen (desensitized with 5% wax) and 25% Al powder; was used in underwater ammunition. [PB Rept 1820, p 40].

Hexamethylenetetramin (Hexamethylenetetramine) (HMTA), called also Hexamin, Methexamine, Aminoform or Ureocapran. See general section.

Hexamethylenetetramine Derivatives (Explosives). To this group belong explosives containing Hexogen (RDX or Cyclotrim) and E-Salt (Cyclotrimethylene Tetramine) described elsewhere. In addition, G.Römer et al investigated two explosives (see Aliphatic Nitramines of WW II) obtained as by-products in the manufacture of Hexogen by the E-Salt and KA-Salt processes.

Both of these substances were claimed to be more powerful explosives than Hexogen.

Reference: G.Römer, PBL Rept 85,160 (1946), p 16.

Hexamethylenetetraamidodiamine (HMTADA) (Hexamethylenetetraamidodiamine). Preparation and properties are given in the general section. The explosive was proposed in 1917 for use as an initiating component for detonators. For instance, the No 8 copper cap might contain 0.1 g of HMTADA and 1 g of TNT.

Reference: C. von Grawert, Ger Pat 274,522 (applied for in 1912, issued in 1914).

Hexamin. One of the German designations for Hexamitridiphenylamine. The same designation was used for Hexamethylenetetramine.

Hexomit, or Hexomit. An explosive used during WW I for cast loading torpedoes, sea mines, and depth charges. It consisted of hexamitridiphenylamine (HNDPPha) 60-70 and TNT 40-50%. Its properties are described in the general section.

After termination of WW I, the Hexomit was used as a component of a commercial explosive known as "Neuclit".

This term Hexomit was also used for the following commercial explosive prep from surplus material of WW I: 60 to 90 parts of HNDPPha, in which might be present up to 40% picric acid, 10 to 40% TNT, TNT, and/or TNN, and 0 to 4% vegetable meal.

Reference: J. Péria Lehuille, Poudres, etc, Paris (1935), pp 437-8.

Note: According to TM 9-1985-2 (1953), p 15, the Hexomit was used in the warfare of Kurt Apperwein (q v).

Hexomit. Same as Hexamin.

Hexamitridiphenylamine. See general section under Diphenylamines.

Hexamitridiphenylamine. Same as Hexa.

Hexa. One of the abbreviations for Hexogen (H) (Cyclotrim or RDX).

Hexa (5-19 and 5-22). German substitute explosive consisting Hexogen (RDX); described under "Ersatzsprengstoffe".

Hexogen or HMDX (also called V-Salt, E-Salt, Z-Salt, SH-Salt and KA-Salt), depending on the method of manufacture. It is described in the general section as Cyclotrim (Cyclotrimethylene Tetramine).

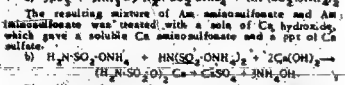
Verfahren zur Herstellung von Ammoniumsalzen
Bemerkung: Der Übersetzungsfehler ist nicht
Dokument, den die D.M.B. Universität
an der Universität Berlin

Verfahren zur Herstellung von Ammoniumsalzen
Bemerkung: Der Übersetzungsfehler ist nicht
Dokument, den die D.M.B. Universität
an der Universität Berlin

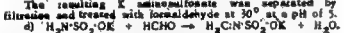
Although Hexogen was known in Germany since 1899 (Hemling, Ger Pat 104 280, 1899), it was not used as an explosive until about 1935 when its manufacture was started using the V-Verfahren described below. Four other methods of manufacture were later introduced and production reached its peak with 100,000 lb produced during the month of June 1945. One of the five methods developed in Germany and described briefly below, the so-called KA-Verfahren proved to be the best because it was the most economical, required less space and equipment and used readily available raw materials.

Following are the German VV II methods of manufacture, arranged in approximate chronological order:

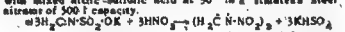
1. V-Verfahren (V-Process), developed in 1935 by Dr. Wolfman of the IG Farbenindustrie, was treated in the literature indicated by the following equations, starting from sulfur dioxide and ammonia:



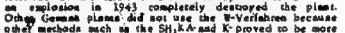
The liberated ammonia was recovered and used in reaction (a). The Ca sulfate was removed by filtration and the Ca amino-sulfonate treated with K sulfate.



The resulting K amino-sulfonate was separated by filtration and treated with formaldehyde at 30° at pH of 5.



The resulting condensation product, K methylhexamino-sulfonate, called Tetra-Sol (white salt), was stirred with mixed nitric-sulfuric acid at 30° in a stainless steel nitron of 500 l capacity.

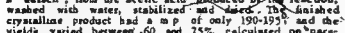


This procedure (which under certain conditions gave yields up to 80% based on the formaldehyde used) was followed at the Krummel Fabrik of Dynamit A-G until explosion of the plant in 1945.

Other German plants did not use the V-Verfahren because other methods such as the SH-K-A and K-Prozess to be more economical.

From a similar method was patented later by R.W. Schlieffert and J. H. Rosen, U.S. Pat. 2,434,730 (1948).

2. E-Verfahren (E-Process), developed between 1935 and 1938 by Drs. Ebeling and Fischer, was based on the reaction of paraformaldehyde with Am. nitrate, dissolved in acetic anhydride, which resulted as a by-product in acetic anhydride.

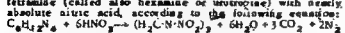


The resulting Cyclohexane was separated by means of a siphon, from the acetic acid produced by the reaction, washed with water, stabilized and dried. The finished crystalline product had a m.p. of only 190-195° and yields varied between 60 and 75% calculated on paraformaldehyde.

The E-Verfahren was used at the Bobingen Fabrik, Dynamit A-G and produced 125 metric tons per month. It was replaced in 1944 by the KA-Verfahren which enabled the production to be doubled with the same equipment.

Note: The Cyclohexane obtained by this method contained the same impurities as described under KA-Verfahren but in larger amount.

3. SM-Verfahren (SM-Process), developed in 1937-1938 by Dr. Scheurer was based on the original method of Henning (1899), which involved direct nitration of hexamethylenetetramine (called also hexamine or urotropine) with acetic anhydride and nitric acid, according to the following equation:



A similar method, was independently developed by Dr. C. H. Hale at Picatinny Arsenal.

The improvement introduced by Dr. Scheurer consisted in carefully controlled heating ("cooling-off") of the contents of the nitator directly after the completion of the reaction.

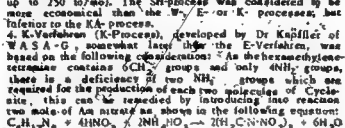
Under these conditions the unstable products formed during the reaction were partly decomposed and partly utilized to cyclonize.

The situation in the SM-process was conducted at -10° using white 99% nitric acid. The purified Cyclohexane had a m.p. between 200° and 107°C.

Was in the original Henning method the yield was very low (about 40% based on $\text{C}_6\text{H}_{12}\text{N}_4$ feed), the improved method was much more economical (yields up to 71.5% were achieved).

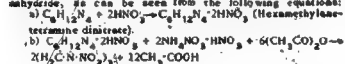
The SM-process was used in at least three plants all of them belonging to the Dynamit A-G's Chiemsee (producing up to 3000 metric tons per month), Döberitz (producing up to 500 t/mo) and Uckermark (producing up to 250 t/mo). The SM-process was considered by the German economy as the most economical since it was inferior to the KA-process.

4. K-Verfahren (K-Process), developed by Dr. Kneffler of W.A.S.A.-G. somewhat later than the E-Verfahren, was based on the following condensation:



Nitric acid of 99% strength was used and was required in larger quantity than for the other methods. This made the recovery of spent acid a very difficult and expensive problem. Only one German plant used this method (Elsäss. Fabrik of W.A.S.A.-G.), producing 200 metric tons per month.

5. KA-Verfahren (KA-Process), developed by Dr. Knöfeler of W.A.S.A.-G. was actually a combination of the K- and E-processes. It consisted in treating the hexamethylenetetramine dinitrate with acid, Am. nitrate in acetic anhydride, as can be seen from the following equation:



In this method, condensed to be one of the most economical, paraformaldehyde was used because all the necessary CH_2 groups were supplied by hexamethylenetetramine. A similar procedure was developed in the U.S.A. by W.E. Bachmann (see page 89).

In the KA-process, as practiced at the Bobingen Fabrik, hexamine was treated with weak nitric acid (35-50%) without heat and the resulting dinitrate was dried in a vacuum oven.

The dry product was dissolved in acetic anhydride, forming a stainless steel vessel equipped with a hydrotrope stirrer, and then acid Am. nitrate (previously prep by treating Am. nitrate with 1 mol of 100% nitric acid) was added. The resulting solid product was separated from acetic acid, then washed with water and dried. The cyclohexane obtained by this method was called KA-Sol. It contained, as impurities, 1 to 2% of HMX (cyclotetramethylenetetramine, called in Germany Detonol), $(\text{H}_2\text{C}-\text{N}-\text{NO}_2)_2$, and a small amount of cyclotrimethylenetetramine, $(\text{CH}_2)_3\text{N}_3\text{O}_3$.

Higher percentages of these impurities were produced when the E-Verfahren was used. Note: The advantage of the KA-process over the E-process was that by using hexamine instead of paraformaldehyde, only half of the amount of water was produced, thus requiring a much smaller amount of acetic anhydride. Hence, it was possible, without increasing the size or amount of equipment, to increase the production of the Bobingen Fabrik, Dynamit A-G from 125 to 250 metric tons per month when the method was changed in 1944 from the SM- to the KA-process.

Yields, when calculated on the basis of formaldehyde (from which the hexamine was produced) were 90-95% for the KA-process, as against 75-75% for the E-process. In the KA-process the production of 100 parts of Cyclohexane required 400 parts of hexamine, 430 parts of nitric acid, and 2400 parts acetic anhydride (of which 19% were recovered as acetic acid).

A recent article of Meyer (Ref. 5) described some German methods of preparation of RDX and their properties as follows: m.p. 201-3, 1.82, explosion

temperature 230°, impact sensitivity by 2 kg weight 40-45 cm, velocity of detonation 8400 m/sec.

Straight Hexogen was used by the Germans as a booster, sub-booster and as a bursting charge in "Villier grenades" and some small caliber shells. It was also used with a small amount of wax, e.g., 3%, as a sub-booster in the African campaign to replace PETN-wax mixtures. With a larger amount of wax, e.g., 10-15%, it was used in 75 mm shells. Hexogen was also used with other proportions of wax as well as with TNT, Al etc. (See Filles Nos 86, 89, 90, 91-95, 92-103, 92-103, 95-97/Ep. 02, 103 (or Trialen 105), 106 (or Trialen 106) and 109 (or Trialen 109), described under Filles.)

References: 1) PB Rept 925 (1945) 2) PB Rept 16,669 (1945) 3) Allied and Enemy Explosives, Aberdeen Proving Ground (1946) 4) A. Stenitzsch, Spreng- und Schietstoffe, Zürich (1948) 5) G. G. 6-69 7) J. Meyer, Explosivstoffe, 1954, No 2/8, pp 83-9 (Über Hexogen, seine Fabrikationsmethoden und Eigenschaften).

Remont. One of the explosives invented by Stenitzsch. See under Swiss Explosives.

Manuscript 75: A plastic explosive, developed during WW II at the Krummel Fabrik of Dynamit A-G. It contained RDX 75, NC 1.2 to 1.4, liquid DNT 20.0 and TNT 3.8 to 3.6%. This mixture was prep by heating the required masses of RDX to 50° in a Vortex-Flinder mixer, and blending it with a small amount of NC. This was followed by the addition of a DNT-TNT mixture and further blending. By using this order of addition, lumping was avoided.

The mixture was put out in cylinders about 220 mm long by 28 mm in diameter. It was difficult to cut direct cap initiation. A booster was provided. It consisted of compressed, palenated PETN pellets about 40 mm long by 21 mm diam and equipped with a detonator well 20 mm deep.

Note: This explosive was developed as a substitute for the plastic explosive, which had RDX plus Ammonium vascello, because the latter component was no longer available in Germany. This vascello, called "Kollon" by Meyer, had much greater adherence than vascello manufactured in other countries.

References: Q.W. Stickle, General Summary of Explosive Development, PB Rept No 325 (1945) Appendix 7 (Riley, Development work on Explosives at Krummel).

Manf., Same as Hex.

High Pressure Pump, See Hochdruckpumpe.

High Speed Tanking (in testing various weapons are described in CIOG Rept 38-47 (1945) and in L.E. Simon, German Research in WW II, J. Riley, N.Y. (1947).

Hochdruckpumpe oder V-3 (High Pressure Pump), called also "Barry Little" (Waldgrave 1 was a constant-pressure gun developed during WW II by Condor, an engineer of the firm Bocking, Stenitzsch, and introduced in the "V" (the "V" being a "V" across the Channel to London. The barrel, called 150 mm (5.9"), was of unalloyed crucible steel (steel) and was of a great many V-shaped sections, each 12 in to 16 in long. With the gun about 150 ft long containing about 28 propellant elements (distributed along the barrel), was reported to achieve a muzzle velocity of about 4200 ft/sec and a range of about 130 km (when using a projectile 8 ft long and weighing 150 lb).

(VERG)

The gun could be used on wooden and concrete targets, and the barrel and the base of the projectile were changed in order after another (if the projectile was damaged by charges between 2 and 5 minutes but because the section was not inserted new 3" projectiles).

References: 1) L.E. Simon, German Research in WW II, J. Riley, N.Y. (1947) 2) W. Waldgrave, V-3, N.Y. (1947) 3) L.E. Simon, German Research in WW II, J. Riley, N.Y. (1947)

Hochgeschwindigkeit (HE). See general.

Maas and Niederdr. Gas, observed to be present in a gun projectile, that it can contain even on

Germany, since 1899 it was not used as an explosive and was abandoned. Four other methods and production reached during the month of development in Germany so-called KA-Verfahren, the most economical, and used readily available

method of manufacture, called: In 1933 by Dr. Wolfram sold on the reaction, starting from sulfur

$H_2S + HN(SO_3)ONH_2$
monosulfonamide and Am
sulfate and a part of Ca
 $NH_4NO_3 + 2Ca(OH)_2$

$SO_2 + 2NH_4OH$
corated and used is
removed by filtration
with a K sulfate.
 $H_2N(SO_3)ONH_2$
was separated by
at 30° at a pH of 5.
 $C_2H_5N(SO_3)OK + H_2O$

product, K methylene
sulfonamide, was stirred
in a stainless steel
 $C_2H_5N(SO_3)OK + 3KHSO_4$

grain conditions, the
granulometry used was
Dynamit A-G until
destroyed the product
K-Verfahren because
it proved to be more

later by R.W. Schlegel
between 1933 and
based on the reaction
by dissolved in acetic
acid.
 $O = (H_2N-C_2H_4-NO_2)_2$

pointed by means of
duced by the reaction,
the finished product
only 190-195° and the
calculated on par-

the Bobagies Fabrik,
nitric tons per month.
reaction which enabled
the same equipment,
this method contained
8 under KA-Verfahren

developed in 1937-1938
the method of chemical
production of hexamethylene
urethane) with nearly
the following reaction:
 $6H_2O + 3CO_2 + 2NH_3$

ately developed by
Dr. Schauer consisted
control of the concentra-
tion of the reaction.

Under these conditions the unstable products formed during the reaction were partly decomposed and partly attached to cyclonite.

The attitude in the SH-process was conducted as follows: using white, 99% nitric acid. The purified Cyclonite had a pH between 200° and 102° C.

While in the original (Hase) method the yield was very low (about 4% based on $C_2H_5NO_2$ used), the improved method was much more economical (yields up to 71.5% were reported).

The SH-process was used in at least three plants all of them belonging to the Dynamit A-G: Christenstam (producing up to 300 metric tons monthly), Oberita (producing up to 200 tons monthly), Uckermünde (producing up to 250 tons monthly). The SH-process was considered to be more economical than the E- or K-processes, but

injection in the K-process. 4. K-Verfahren (K-Process), developed by Dr. Kapffler of W.A.S.-G., somewhat later than the E-Verfahren, was based on the following considerations: 2. In hexamethylenetetramine contains CH_2 groups and only CHN groups, there is a deficiency of two NH_2 groups which are required for the production of each two molecules of Cyclonite, this can be remedied by introducing into reaction two moles of An nitrate as shown in the following equation:

$6HNO_3 + 2NH_4NO_3 \rightarrow 2H_2N(C_2H_4NO_2)_2 + 5H_2O$
Calcium acid of 95% strength was used and was required in larger quantity than for the other methods. This made the recovery of spent acid a very difficult and expensive operation. Only one German plant used this method (Fahrik of W.A.S.-G.), producing 200 metric tons per month.

5. KA-Verfahren (KA-Process), developed by Dr. Kapffler of W.A.S.-G. was actually a combination of parts of the K- and E-Processes. It consisted in treating the hexamethylenetetramine distillate with acid. An nitrate in acetic anhydride, it can be seen from the following equation:

a) $C_6H_{12}N_4 + 2HNO_3 \rightarrow C_6H_{12}N_4 \cdot 2HNO_3$ (Hexamethylenetetramine dinitrate).
b) $C_6H_{12}N_4 \cdot 2HNO_3 + 2NH_4NO_3 \cdot HNO_3 + 6CH_3CO_2O \rightarrow 2H_2N(C_2H_4NO_2)_2 + 12CH_3COOH$

In this method, considered to be one of the most economical, formaldehyde was produced by this method (Necessary CH_2 groups were supplied by hexamethylenetetramine). A similar procedure was developed in the U.S.A. by Dr. E. Schuchman (See also Chapter 10).

In the K-Process, as practiced at the Bobagies Fabrik, hexamine was treated with weak nitric acid (55-50% nitrous acid) and the resulting dinitrate (called in Germany Hexedit) was dried. The dry product was dissolved in acetic acid (hydro-type nitrate) and then acid An nitrate (previously prepared by treating An nitrate with nitric acid) was added. The resulting solid product was separated from acetic acid, then washed with water and dried. The cyclonite obtained by this method was called KA-Salt. It contained, as impurities, 1 to 2% of MMX (cyclohexamethyleneurethane, called in Germany Oetogen), $(H_2N-C_2H_4-NO_2)_2$ and a small amount of cyclonitrimide (dinitrodioxyethylene), $(CH_2)_6N_4(NO_2)_2OCH_3$. Higher percentages of these impurities were produced when the KA-process was used.

Note: The advantage of the KA-process over the E-process was that by using hexamine instead of paraformaldehyde only half of the amount of water was produced, thus requiring a much smaller amount of acetic anhydride. Hence, it was possible without increasing the size or amount of equipment, to increase the production of the Bobagies Fabrik, Dynamit A-G, from 125 to 250 metric tons per month when the method was changed in 1944 from the E- to the KA-process.

In fields, when calculated on the basis of formaldehyde (from which the hexamine was produced) were 90-92% for the KA-process, as against 73-75% in the E-process. In the KA-process the production of 100 parts of Cyclonite required 40p of hexamine, 43p of An nitrate and 240p of acetic anhydride (of which 159p were recovered as acetic acid).

A recent article of Meyer (Ref 3) described some German methods of preparation of RDX and lists its properties as follows: up 201-5, 1.82, explosion

temperature 230°, impact sensitivity with 2 kg. weight 40-45 cm, velocity of detonation 3400 m/sec.

Straight Hexogen was used by the Germans as a booster, booster-booster and as a bursting charge in "Riffengroßer" and some small caliber shells. It was also used with a small amount of wax, c.a. 3%, as a sub-booster in the African campaign to replace PETN-wax mixtures. With a larger amount of wax, c.a. 10-15%, it was used in 75 mm shells. Hexogen was also used with other proportions of wax as well as TNT, Al etc. [See Fillers Nos 86, 87, 89, 90, 91-95, 92-100, 3, 52-63, 93-94/E 02; 105 (on Trialen 105), 106 (on Trialen 106) and 109 (on Trialen 109), described under Fillers].

References:
1) P.B. Rept 925 (1945) 2) P.B. Rept 16,669 (1945) 3) Allied and Enemy Explosives, Aberdeen Proving Ground (1946) 4) A. Schuchman, Spang and Schienkowitz, Zürich (1946), pp. 68-69 5) J. Meyer, Explosivstoffe, 1954, No 2/8, pp. 83-5 (Ubers. Nitrogen, and Fertilizationsmethoden und Eigenschaften).

Hexamin. One of the explosives invented by Schuchman. See under Swiss Explosives.

Hexamit 75. A plastic explosive, developed during WW II in the Krimmel Factory of Dynamit A-G. It contained RDX 75, NC 1.2 to 1.4, liquid DNT 20.0 and TNT 5.8 to 3.6%. This mixture was prepared by heating the required amount of RDX to 90° in a Werner-Pfeiderer mixer, and blending it with a small amount of NC. This was followed by the addition of a DNT-TNT mixture and further blending. By using this order of addition, lumping was avoided.

The mixture was put out in cylinders about 220 mm long by 28 mm in diameter. Due to difficulty with direct extrusion, a booster was provided. It consisted of compressed, phlegmatized PETN pellets about 40 mm long by 21 mm diam and equipped with a detonator well 20 mm deep.

Note: This explosive was developed as a substitute for RDX, which was used RDX plus American vaseline, because the latter component was no longer available in Germany. This vaseline, called "long flosser" by Meyer, had much greater advance than vaseline manufactured in other countries.

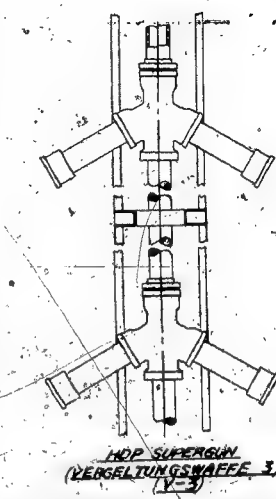
Reference: O.W. Sackel, General Summary of Explosive Plants, P.B. Rept No 525 (1945) Appendix 7 (R. Meyer, Development 60th on Explosives at Krimmel).

Hexyl. Same as Hexan.

High Pressure Pump. See Hochdruckpumpe.

High Speed Tooling (for testing various weapons are described in CLOS Rept 22-47 (1945) and in L.E.Slim, German Research in WW II, 1. Bailey, N.Y. (1947).

Hochdruckpumpe oder V-3 (High Pressure Pump, called also "Long Lister" or "Marsden") was a continuous-pressure gun developed during WW II by Condor, an engineer of the firm Beckling, Starckebach, and intended to fire the 400 mm (Hase) Projector (Rept 22-47) across the Channel to London. The barrel, caliber 150 mm (5.9"), was of unalloyed crucible cast steel made up of a great many tapered sections; each 12 to 16 ft long. The gun about 450 ft long, containing about 28 propellant chambers (distributed along the barrel). It was expected to achieve a muzzle velocity of about 4500 ft/sec with a weight of about 150 lb (when using a projectile 8 ft long and weighing 150 lb).



The gun could fire on the ground without any carriage on wooden and concrete blocks angled at a 45° angle. The forward end, where projectiles were inserted in the barrel and the base propellant charge electrically ignited. As the projectile passed the separate V-pieces, additional propellant charges in the side arms were electrically ignited one after another (in pairs) thus accelerating the velocity of the projectile as it progressed along the gun barrel.

For servicing (reloading the V-pieces with propellant charges between the rounds), the gun required a great many soldiers. It was planned to fire one round per gun every 3 minutes but this rate could not always be achieved because the sections after exploded and it was necessary to insert new V-pieces.

References:
1) L.E.Slim, German Research in World War II, Viking, N.Y. (1947), pp. 12-13 2) W. Lohmeyer, V-2, Viking, N.Y. (1954), p. 247 3) A.L. Sprain and H.H. Ballock of Picatinny Arsenal; private communication.

Nochmischelaststoffe oder Blittrstoffe (High Explosives) (HE). See general section.

Mech. and Mischdruckmischungen (High and Low Pressure Gun, abbreviated to H/L-Gun) (Cimon & Meyers, in French). It has been known for a long time that the lower the peak pressure in a gun the thinner may be the walls of the projectile. This means that for a given total weight of a projectile, that used in a gun with lower peak pressure can contain more explosive and do more damage to a target.

of shaped charges
es not depend upon
in the amount of the
low pressure in a
e, should be made
case larger. Such
suitable because the
it burned irregularly
er). Also, the initial
ound to round which
achieved.

13 when Dr Hermann
-Borsig A-G con-
Antitank Gun). The
ch "canon antichar
Touchard (Ref. 3).
hereafter" invented
be considered as the
ess guns.

a comparatively thin
and was 34 calibers
meter (105 mm)
dual type, 81 mm
le and weighed 314 kg.
epacite-loading am-
tridge (120 mm long
ided the propellant.
of a disc provided
in diameter). When
the gases developed
cm, but the pressure
kg/cm² because the
passing through the

pressure inside the
e in the bore could
the size or number
In order to protect
spilling and from
was covered with a

ere worked out by
rmer in England
end of WW II the
light antitank guns:
PAW 1000, but does
the 8.8 cm W7 gun,
"snipe".

(1948) 2) J. Corner,
Guns, J. Wiley, N.Y.
L. Touchard, Mém
22, 219-36 & 245-78

Considerable work
WW II on the devel-
most prominent con-
3 of Krummel's Pat-
eguns developed as

age weighing 3 kg;
240 mm
atrons, Panzerfaust,
best explosives the

shaped charges were RDX-TNB and next, RDX-TNT mixtures.
Sustitut. PETN for RDX lead to a decrease in efficiency.
The addition of aluminum powder was desirable but not
in large quantities.

Krummel was not the only place where work on shaped
charges was conducted. Elsewhere the Germans developed
a shaped charge "well" which was shot from an 80 mm mortar
called "Panzerwulkan", and the warheads for several
guided missiles.

Historical Discovery of the hollow (shaped) charge
(HoC) effect is usually attributed to C.E. Munroe (U.S.A.)
who described the effect in the Amer. J. Sci. 36, 1888. It
was claimed by H. Schardin that Max von Forster of
Germany had in 1883 already shown that bare hollow
charges gave an enhanced effect along the axis of the
charge. The first practical application of the HoC effect for
demolition charges, sea mines, torpedoes, projectiles
etc., was presented in 1910 by E. Neumann & the head
(Kilisch-Anhaltische Schmelzwerk A.G. (R.P. Ann. v. 1629).
Neumann's work is described in S.S. 9, 183 (1914).
S.S. 9, 183 (1914). Important work on this subject
of the HoC effect was done prior and during WW II. By
H. Schardin et al. in Berlin. Some work was also carried
out by A. Stettbacher of Switzerland during this period.

Note: According to A.J. Dene, Ordnance Sergeant, October
1945, p. 3-13, hollow (shaped) charge ammunition was
used by the Germans in many 75 mm caliber weapons.
There were at least four types of such projectiles: H1,
H1/A, H1/B and H1/C. Most of these projectiles are listed
in the disciplinary under Granate and are briefly described
in TM 3-178-53 (1953). Some projectiles of calibers 88 mm,
100 mm, 105 mm and 150 mm also had shaped charges.
The enclosed drawings represent some typical German
hollow charges. (See next page).

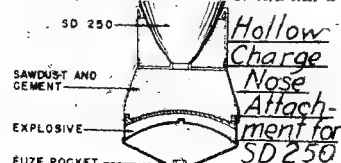
References:
1) A. Stettbacher, Nitrocellulose 8, 83-84 (1937)
2) O.W. Stickland et al, PB Rept 925, Appendix 3, p. 46
and Appendix 7
3) L.E. Simon, German Research in WW II, Wiley, N.Y.
(1947), pp. 118-120, 188
4) A. Stettbacher, Spreng- und Schießstoffe, Rascher, Zürich
(1948), pp. 133-34
5) H.L. Potter et al, CIOS Report 33-27 (1945). This
report is classified and information contained therein
has not been used for this document.

(See also Shaped Charge in the General Section)

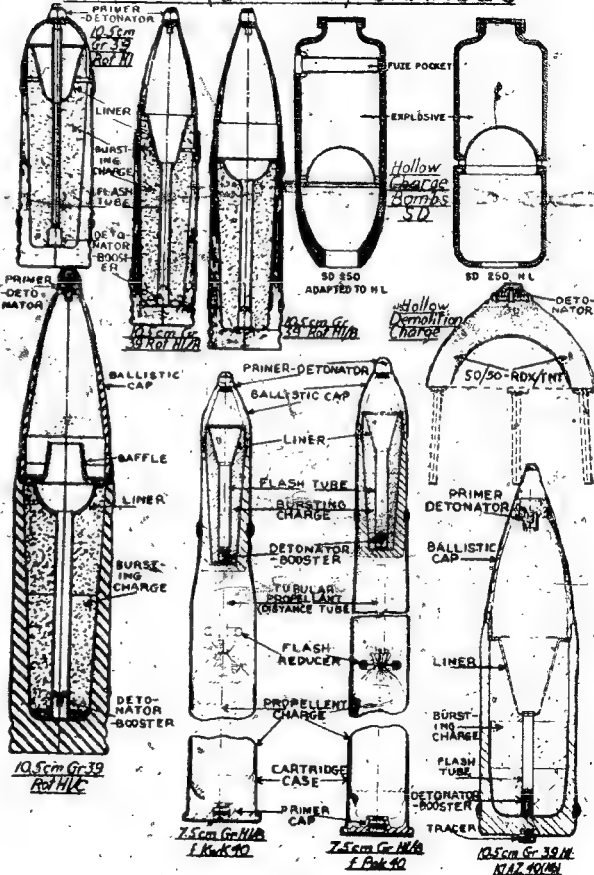
Haka (Hochkonzentriert, a highly-concentrated) Process
for the manufacture of 98-99% nitric acid developed during
WW II was used in several German plants. In this process,
the concentration of the weak acid (50%) was effected by
mixing it with liquid nitrogen tetroxide (N₂O₄) and adding
the necessary extra oxygen under 30 atm pressure in an
autoclave.

Description of this method as practiced by the IG
Faserland A-G subsidiary, the Virschafstliche Forschungsgesellschaft,
mbH (VfG), Embsay, Kz. Lüneburg is given
in the following BODS Final Report: 1232 (1947), pp. 15-16
and 1442 (1947), pp. 84-86.
Hollow Charge. See Hollowbombs.

Hollow Charge Nose Attachment for AP Bombs. In order to



HOLLOW (SHAPED) CHARGES



permit greater penetrating power from low altitudes some German 250 kg AP bombs had a hollow charge (weighing about 4 kg) attached to the nose. This charge was detonated by its own nose fuse as soon as it hit the armor. The explosion of the HoC produced a hole in the armor (as deep as 7 cm) which permitted the AP bomb to enter inside the target. The AP bomb being provided with a short delay fuse did not explode until it was inside the target. In order to protect the bomb from pressure detonation the space between the HoC and the nose of the bomb was filled with sawdust and cement.

Reference: TM 9-1985-2 (1953), p 3.
Holsapfen (Wood Spins). See Methanol in general section.

Holzwoll (Wood Wool). See Wood Fluff in the general section.
Holzleiste 42-See under Landminen and also on p 263 of TM 9-1985-2 (1953).

Holzsapfen (Wood Pich). See general section.

Holzschneeflocke (Wood Pulp). See general section.

Holzstiel (Wood Tac). See general section under Tac.

Holzstoffsack (Wood Cellulose or Chemical Wood Pulp). See general section.

Naming Guidance Systems for Missiles, such as Acoustic, Radar and Infrared are briefly described under Guidance Systems for Missiles.

Hawitze (Haußitz). See under Weapons.

Ha 117 (Henschel-117), also known as Schmetterling (Butterfly), was a rocket propelled, radio controlled, missile for use against bomber formations. Some versions were for ground use and some for air-to-air. It used liquid fuel called Tonka and as oxygen carrier called Salbei.
[TM 9-1985-2 (1953), pp 196-201].

Ha 293 (Henschel 293) was a radio-controlled missile released and directed to the target from an aircraft. The model fully developed and used was the Ha 293 A-1. Other models such as Ha 293 A-2, Ha 293 B, Ha 293 C, Ha 293 D, etc were not fully developed. [TM 9-1985-2 (1953), pp 201-3].

Ha 296 (Henschel 296) was a rocket-propelled, radio-controlled missile designed primarily as an air-to-air weapon to be carried on fighter aircraft as well as the bomber types. There were several versions but the basic type was called Ha 296 V-2. It used a solid propellant.
[TM 9-1985-2 (1953), pp 205-5].

HTA. An abbreviation for mixtures of RDX (Hexogen), TNT (Trinit) and Al (aluminum), such as in the proportions 40/40/20. [See also RBL Rept No 83,160 (1946), p 15].

Hydrazin Propellanten, presented in 1895, were prepd by mixing AC (gelatinized by means of 2-3% solution of K-sulphuric ether-alcohol) with small quantities of nitrosophthal, nitrobenzoesäure, or nitrobenzol. For instance, a propellant used for military purpose contained 4 to 5% of nitrosophthal, (David, Dictionnaire, Paris (1902) p 478).

Hummel (Bumble Bee). Nickname for a self-propelled mount consisting of 150 mm medium howitzer on the chassis of a PzKpfw III/IV tank. (See also under Panzer).

Hydrazine. Hydrazine is described in the general section. Its manufacture in Germany at IG Farbenindustrie Planten at Gersthofen, Leverkusen, Ludwigshafen, is described in BIOS-Final Report (1946).

Hydrocellulose. Hydrocellulose (Hydrocellulose).

Described in the general section. It was reported that the Germans used it in some rocket propellants, presumably to improve the burning characteristics. For instance the so-called Ammonipulver contained 5% hydrazine and the EP (Eisstaupulver) contained about 3%. Hydrocellulose was also used in some rocket propellants to increase the rate of burning at low temperatures. (See also under Propellant). Reference: CIOS Report 31-68 (1945), pp 6-7.

Hydrogen Peroxide (Wasserstoffsuperoxyd). Its preparation and properties are described in the general section under Peroxides. It was used in liquid rocket propellants and in a special tubing designed for submarines, by Walter. Several German methods of manufacture are described in the following References:

- 1) B.E.A. Vigor et al. Hydrogen Peroxide Production by Electrolysis of 35 Per Cent Solutions (Deutsche Gold und Silber Anstalt), BIOS Final Report 683 (1945).
- 2) V.W. Slater et al. The Anthraquinone Autoxidation Process for the Production of Hydrogen Peroxide, CIOS Report 31-45 (1945).
- 3) J.M. Aulay, Hydrogen Peroxide Manufactured by All-Liquid Process From Ammonium Persulfate, (NH₄)₂S₂O₈, CIOS Rept 33-43 (1945).
- 4) J.M. Aulay, Direct Synthesis of Hydrogen Peroxide by Electrolytic Discharge, CIOS Rept 33-44 (1945).

[See also T-Stoff, Rocket Propellants, Liquid and U-Boat (Unterseeboot) of Walter].

Hygroskopische oder Feuchtigkeits (Hygroscopicity, Humidity or Moisture). Methods of determination are given in the general section.

Igniter (Zünder). The following igniters are briefly described or listed in Reils 1, 2 & 3:

- a) A. Section (Pul) Type (Brennzünder).
- b) BZ 24, with delay pellets, was used in stick grenades (Reif, p 83,13 & 3, p 283).
- c) NBZ 34, with delay pellets was used in smoke grenades (1, p 83,13 & 3, p 283).
- d) BZ 35, with pellet, was used with egg grenades, having stick grenades and message box (Reif, p 1, p 83,12 & 3, p 284).
- e) BZ 39, used in smoke hand grenades (3, p 285).
- f) Zische ANZ 29, used to ignite safety fuse of detonators, to set booby traps, to ignite safety fuses for some demolition charges, to ignite some smoke candles and to booby-trap some Teller mines and grenades (1, p 83,10 & 3, p 285).
- g) Zische ANZ 39, used for the same purposes as above (1, p 83,11 & 3, p 289).
- h) BZ 42, delay 4% sec; same as indicated (1, p 165).
- i) Pressure Type (Druckzünder).
- a) DZ 35(A), used in heavy antitank mine and some prepared charges (1, p 83,03 & 3, p 295).
- b) DZ 35(B), used in some booby traps and prepared mines (1, p 83,03 & 3, p 295).
- c) Hebelzündler (Lever Igniter), also called Schube Igniter, consisted of an inverted L-shaped tube, the vertical arm of which was screwed into a mine. The horizontal arm contained the percussion cap, striker spring and striker retaining pin. On top of the arm was attached a lug, on actuating lever (consisting of a hollow metal-tripping piece pivoted on a rivet) and a safety pin. After removing the pin, the downward pressure (as little as 40 lb) on the actuating lever forced out the striker retaining pin, which caused the striker to fire. The actuating lever igniter was used in Clearance Mines (see also under the Buck Igniter) and in self-propelled mines (1, p 83,14 & 3, p 296).
- d) RZ 42, used in some improvised mines (1, p 83,03 & 3, p 297).
- e) Weisensche Igniter consisted of a spring loaded striker bolt at the top of which was a pressure head. The bolt was held against the spring by a safety device consisting of a small pair of tongs. After removing the tongs, pressure or a blow on the pressure head shattered the glass rod, thus allowing the spring to drive the

Wissenschaft für anorganische Physik Groß
Bauwerke - Untersuchungen - vom Institut
Düsseldorf den 28.11.83. Universität
Sigmund

erf. für dienstliche Zwecke der Kampfmittelbeseti-
gung, Weitergabe an Dritte nur mit Zustimmung des IM TM

power from low altitudes some had a hollow charge (weighing nose. This charge was detonated soon as it hit the armor. The fused a hole in the armor (as the AP bomb to enter inside tank provided with a short delay it was inside the target. In from premature detonation of and the nose of the bomb was not).

e Methanol in general section.

pod. Flow in the general section.

Indpinner and also on p 263 of

General section.

See general section.

સામે સરકારના બંધારા દ્વારા,

of Chemical Wood Pulp). See

for Missiles, such as Acoustic,
elfy described under Guidance

2. **ଉତ୍ତରାଧିକାରୀ**

known as Schmetterling (Butterfly), radio controlled, missile functions. Some versions were for air-to-air. It used liquid fuel carrier called Salbei.

as a radio-controlled missile
be target from an aircraft. The
used was the Ha 293 A-1. Other
Ha 293 B, Ha 293 C, Ha 293 D,
TM 9-985-2 (1953), pp 201-2

is a rocket-propelled, radio-controlled aircraft as well as the Bomber versions but the basic type used a solid propellant.

mixture of RDX" (Hexogen),
am), such as in the proportions
Rept No 85,160 (1946), p 13].

in 1895, were prepd. by mixing
of 2-3% soln of K xanthogenate
all quantities of nitronaphthol,
r. For instance, a propellant
contained 4 to 5% of nitronaphthol,
(1902) p 378.

name for a self-propelled mount-
ing Howitzer on the chassis of
(also under Panzer).

cribed in the general section.
any at the IG Farbenwerke
Kruen, Ludwigshafen
1. 1. 1946.

Hydrocellulose (Hydrocellulose).

Described in the general section. It was reported that the Germans used it in some rocket propellants, presumably to improve the burning characteristics. For instance the so-called Ammonpulver contained 3% hydrocellulose and the EP (Einheitspulver) contained about 3%. Hydrocellulose was also used in some rocket propellants to increase the rate of burning at low temperature. (See Standard Propellant). Reference: CIOS Report 31-68 (1945), pp 6-7.

Hydrogen Peroxide (Wassersstoffsuperoxyd), its preparation and properties are described in the general section¹ under Peroxides. It was used in liquid-rocket propellants and in a special turbine designed for submarines by Walter. Several German methods of manufacture are described in the following references:

2) V.W.Slater et al., The Anthraquinone Autoxidation Process for the Production of Hydrogen Peroxide, CIOG-Report 31-15 (1945)

3. J. McAulay, Hydrogen Peroxide Manufactured by All-liquid Process From Ammonium Persulfate, (NH₄)₂S₂O₈, C. S. Rep. 33-43 (1945).

4) J. McCauley, Direct Synthesis of Hydrogen Peroxide by Electric Discharge, CIOB Rep 33-44 (1945).

[See also T-Stoff, Rocket Propellants, Liquid and U-Boat, (Unterseeboot) of Walter].

Hygroscopicität oder Feuchtigkeits (Hygroscopicity, Humidity of Moisture). Methods of determination are given in the general section.

igniter (Zünder). The following igniters are hereby described or listed in Refs 1, 2 & 3:

A. Friction (Pull) Type (Brennender).

b) NbBZ 38, with delay pellets, was used in stick grenades (Ref 1, p 83, 23 & 3, p 283)

c) HZE, with pellet, was used with egg grenades

d) BZ 39, used in smoke head grenades (3, p 285)

e) 2d Sch. ANZ 29, used to ignite safety fuses of detonators; to set booby traps, to ignite safety fuses for some demolition charges.

for some demolition charges, to ignite some smoke
candles and to booby-trap some Teller mines and
grenades. (11 p 83, 10 & 3, p 285)

1) ZuSchnANZ 39, used for the same purposes as above
(1, p 83.11 & 3, p 285)
2) BZ 42, delay 4 1/2 sec; uncorrected indicated (1, p 16)

B. Pressure Type (Druckstiller).

a) DZ 35(A), used in heavy antitank mines and some prepared charges (1, p 83.03 & 3, p 295)

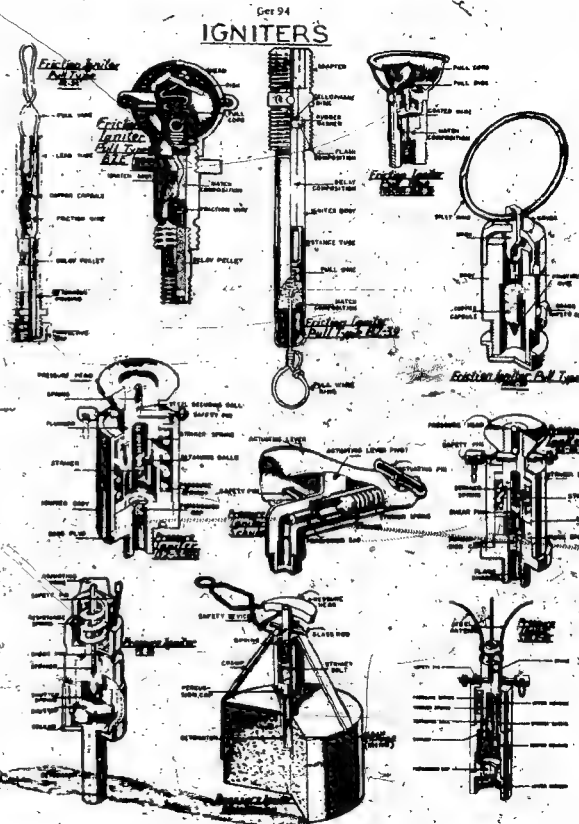
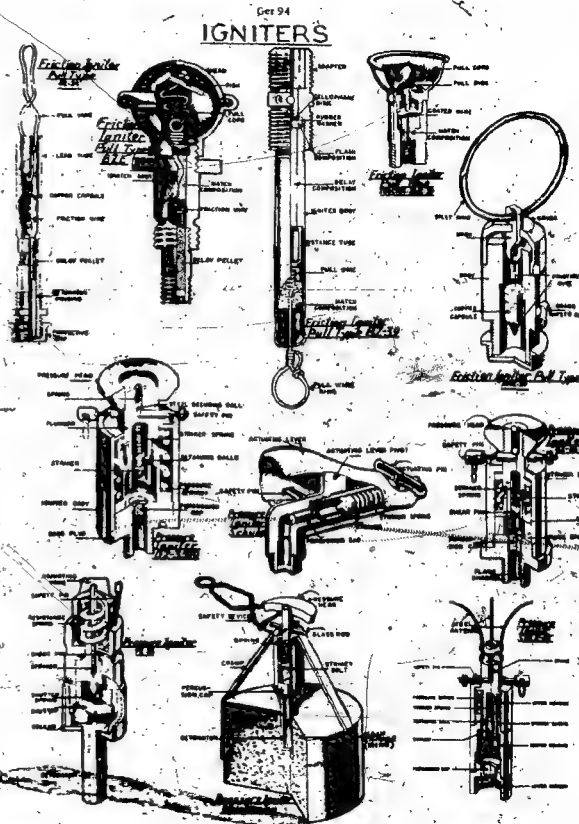
c) Hebelzündler (Lever Igniter), also called Schuko

hammer, consisted of an inverted L-shaped tube, the vertical arm of which was screwed into a mine. The horizontal arm contained the percussion cap and striker.

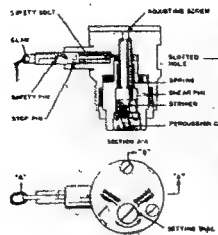
striker spring and striker retaining pin. On top of the arm was attached a lug, an actuating lever (consisting of a hollow metal tripping piece pivoted on a rivet), and a safety pin. After removing the pin, the downward pressure (as little as 40 lb) on the pressure lever forced out the striker retaining pin, thus releasing the striker to fire the device. A Buck igniter was used in Glasgow. A 40 lb downward force on the Buck igniter and in some boats (p. 1, p. 83, 14 & 3, p. 296).

(1) Weapons "limiter" consisted of a spring loaded

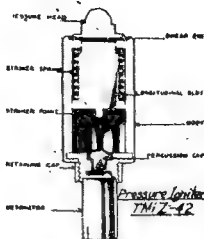
The bolt was held against the spring by a safety device



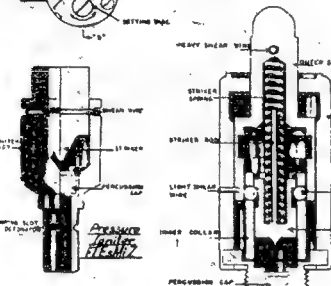
Ger 95 IGNITERS



Pressure Igniter
 TMI-Z-35



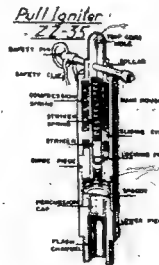
Pressure Igniter
 TMI-Z-42



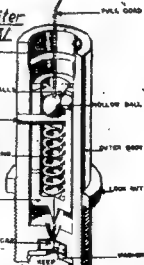
Pressure Igniter
 TMI-Z-500a



Pressure Igniter
 TMI-Z-500b



Pull Igniter
 Z-2-35



Pull Igniter
 Z-2-35

striker against the percussion cap etc. The is designed for use as a push igniter in improv or as an impact igniter for HE charges when u assault (3, p 280)

g) TMI-Z 35 designed for use in Schützeng called Bounding Mine (3, p 293)

h) TMI-Z 35 (Tellerminen-Zunder 35), used in (3, p 301)

i) TMI-Z 42, used TMI-35 (steel), TMI 42 and also called Pilzmine (Mushroom Mine) (3, p 303)

j) TMI 43, used as above (3, p 304)

k) F15TMI-Z, used in Flaschenmine 42 (Ant glass bottle mine) (3, p 307)

l) MZ 550C, an igniter made in Germany in the British Antarctic Mine 330 (3, p 305)

m) Topfminen-Zunder (Pot Mine Fuse) consists hollow, cylindrical, glass body into which a solid pressure head, inside the cylinder were two glass ampoules containing liquids which ignited the explosive train of the Pot Mine (T A pressure of about 150 kg was sufficient the ampoules (3, p 306)

C. Pull Type (Zündarten)

a) Z-2 35, used in S-Mines, some prepared booby traps employing trip wire) and for booby traps (3, p 280)

b) Type 31 designed for use in antiperson and booby traps (3, p 280)

D. Pull and Shear Type (Zug- und Zerschneide) also called Pull and Tension Wire Igniter, Z-2 Z-2 35, consisted of brass cap containing cushion cap, striker, striker spring (located sliding cylinder and held on top by a plunger), compression spring, a retaining (locking) p safety pin. The top of plunger was connected trip wire held under tension. The igniter either by pulling on the trip wire or by loosening or breaking it. In the first case the trip was the plunger to be pulled upward against the r of the outer spring. This permitted the two pins to be forced outward into the upper open thus freeing the striker. In the second case, or cutting of the trip wire allowed the outer spring to force the sliding cylinder downward, permitted the locking pins to be forced out the lower open space, thus freeing the stri Igniter was used with S-Mines-booby traps armed charges (3, p 83.05 & 3, p 293)

E. Percussion Type (Schlagzunder oder Aufschlag) a) Schlagzunder 35 was a modified version of uses not indicated (2, p 163)

b) Safety Fuse Igniter consisted of a cylindrical body containing a spring-loaded striker held in by a friction fit of the 2 type with a cap w attached a large steel ring. A strong pull on detached the striker release plate from the su permitting the spring to drive the striker into cushion cap. The device was used to initiate t, r (3, p 287)

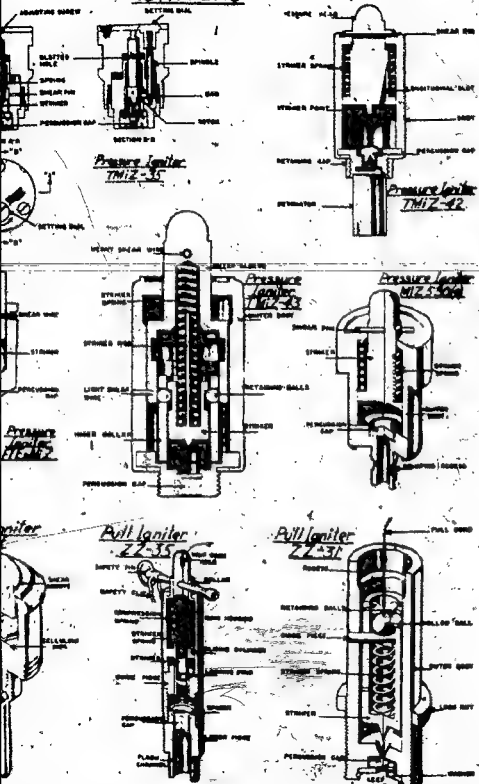
c) Type 2 (Pull Percussion) Igniter was des use with the new type parachute antiperson but was suitable for use with mines and booby For operation, a sharp pull on the split ring on the striker release plate to be drawn from the ign thus releasing the striker spring, which w tension (3, p 280)

d) Aufschlagzunder 355(h) for use in Dutch Mine 355 (3, p 164)

F. Pull and Pressure Type (Zug- und Druckzünd) a) Z-2 29 Igniter, used in the assembly of antivehicle or antipersonnel mines, could be either by pull on a trip wire attached to the pull pin, or by pressure against the set (3, p 292)

b) Z-2 42, consisted of a bakelite cylinder containing a percussion cap, a striker retaining and a striker spring held under tension by wire loop. Pulling on the trip wire attached release pin withdrew the pin thus allowing t to hit the percussion cap. The igniter could operated by attaching a trip wire under stress to the end hole in the striker and carefully the release pin. This igniter was designed f

IGNITERS



striker against the percussion cap etc. The igniter was designed for use as a push igniter in improvised mines, or as an impact igniter for HE charges when used in a usual (3, p 290).

a) SMZ 35 designed for use in Schützengraben, also called Bounding Mine (3, p 299).

b) TMI-2 35 (Eisenkammer 35), used in T-MI 35 (3, p 301).

c) TMI-2 42, used TMI-35 (steel), TMI-42 and TMI 43, used as above (3, p 334).

d) HE-MZ used in Flammenmine '42 (Antipersonnel glass bottle mine) (3, p 307).

e) MZ 350 (an igniter made in Germany for use in the British Detonac Mine 310 (3, p 305)).

f) Topfenkammer (Pot Mine Fuse) consisted of a hollow, cylindrical glass body into which fitted a solid pressure head, inside the cylinder were located two glass ampoules containing liquids which on mixing ignited the explosive tuff of the Pot Mine (Topfenmine). A pressure of about 150 kg was sufficient to crush the ampoules (3, p 306).

g) Pull Type (Zugmine), some prepared charges, booby traps employing trip wires) and for booby trapping of Teller mines (3, p 85-86 & 3, p 282).

h) Type 31 designed for use in antipersonnel mines and booby traps (3, p 280).

i) Pull and Shoe Type (Zug und Zerschneidezünder), also called Pull and Tension Wire Igniter, such as Zuz 35, consisted of a brass case containing a percussion cap, electric striker spring (located in a sliding cylinder and held on top by a plunger), an outer compression spring, a retaining (locking) pin and a safety pin. The top of plunger was connected to a trip wire held under tension. The igniter was fired either by pulling on the trip wire or by loosening (cutting or breaking it). In the first case the trip wire caused the plunger to be pulled upward against the resistance of the outer spring. This permitted the two locking pins to be forced outward into the upper open space, thus forcing the striker. In the second case, breaking or cutting of the trip wire allowed the outer (compression) spring to force the sliding cylinder downwards. This remained the locking pins to be forced outward into the lower open space, thus freeing the strikers. This igniter was used with Mines, booby traps and prepared charges. (3, p 85-86 & 3, p 282).

j) Perforation Type (Schlaglöcher oder Aufschlagzünder).

k) Schlaglöcher 35 was a modified version of Zuz 35, same not indicated (3, p 103).

l) Safety Fuse Igniter consisted of a cylindrical brass body containing a spring-loaded striker held in position by friction fit of the type with a cap to which was attached a large steel ring. A strong pull on the ring detached the striker release plate from the striker thus allowing the spring to drive the striker into the percussion cap. The device was used to ignite a safety fuse (3, p 287).

m) Pull Perforation Igniter was designed for use with the new type antipersonnel mine. The device was suitable for use with mines and booby traps. For operation, a sharp pull on the split ring caused the striker release plate to be drawn from the igniter body thus allowing the striker spring, which was under tension (3, p 288).

n) Aufschlagzünder 3550 (for use in Dutch Antitank Mine 355 (3, p 164)).

o) Pull and Pressure Type (Zug und Druckzünder).

p) Zuz 29 Igniter, used in the assembly of antitank, antivehicle or antipersonnel mines, could be operated either by pull on a trip wire attached to the loop of the pull pin, or by pressure against the setting head (3, p 329).

q) Zuz 42, consisted of a bakelite cylindrical casing containing a percussion cap, a striker retaining washer and a striker spring held under tension by the trip wire loop. Pulling on the trip wire attached to the release pin on the striker pulled the lever into position to hit the percussion cap. The igniter could also be operated by attaching a trip wire under strong tension to the end hole, causing the cap to be pulled against the release pin. This igniter was designed for use in

Stock mine and booby traps (3, p 83-86 & 3, p 193). Note: This igniter is listed in Ref 1 as "Pull" Type, whereas Ref 3 lists it as "Pressure and Pull" Type.

r) SMZ 44, developed for use in S-Mine 14 and in some improvised mines, consisted of a steel cylindrical case containing a percussion cap, striker and spring, winged detents, to which two trip wires were attached.

s) The striker was retained in a cocked position by two (mounted on the case) and a safety pin. After arming the device (by withdrawing the safety pin), a pressure of 21 lb or a pull of 14 lb on the winged detents opened them sufficiently to release the striker.

t) Electric Type (Elektrischer Zünder), ESMZ 40 provided with an electric, Goch funnel-shaped housing, a release plunger, a glass ampoule and two electrodes.

u) In order to enlarge the igniter area for one mine, usually an S-Mine, eighteen of these were connected in parallel, nine igniters in each chain, and spiked in the ground around the mine. The string were connected by means of wires to two plugs fitted into sockets of the electric bridge (aluminum wire), surrounded with a glass composition.

v) The electric bridge was screwed on to the mine Pressure on the prongs of any of the 18 igniters, depressed the release plunger and liberated the two steel locking balls in the head of the strikers. This caused the spring to drive the striker into the glass ampoules. The liberated electrolyte set up a current between the electrodes and the current was transmitted to the bridge wire. The heat of the wire fired the flash composition and finally exploded the HE charge of the mine (1, p 81-83 & 3, p 100-1).

w) Chemical Igniter (Chemischer Zünder).

x) "Buck" Igniter (Chemical Cuck-Accumulated Type) used with the antipersonnel "Pot" mine, consisted of a thin aluminum foil drum containing a glass ampoule with sulfuric acid surrounded with a white, powdered flash composition. The drum was secured by crimping to the brass base. When pressure was applied, the foil drum was dented, the ampoule broken and the acid mixed with the flash composition. This resulted in a chemical reaction which ignited the mine (3, p 108-9).

y) Zuz 419 (Chemisch-mechanischer Zünder), used for delayed action devices consisting of a bakelite housing containing a glass ampoule and other lenses shown on the drawing. When the ampoule was broken by pressure (caused by the impact through perforation in the plastic lid into the reaction chamber (plastic cylinder) where the material was located.

z) As soon as the rod was sufficiently rod weakened and broke, the spring was released thus allowing the striker to hit the percussion cap. The striker was held by the detonator, booster and the main HE charge (3, p 313-14).

aa) All Explosive Pressure Release Device, designed for use as a booby trap, was also suitable as an igniter in mines and other items. The body of the device consisted of two oblong blocks of bakelite (3, p 109-10).

bb) (believed to be Nipolit), held together by two hollow head bolts. The inner surfaces of both blocks were provided with molded recesses to retain the metal striker mechanism. For operation, the device was placed under the object to be booby-trapped so that as the object was lifted the striker retaining arm of the device pivoted upwards, thus releasing the striker which fired the percussion cap (3, p 109-10).

cc) Lamp-Delay Clockwork Igniter.

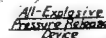
dd) 21-Day Delay Igniter was used in conjunction with large electric demolition charges (3, p 109-10).

ee) 21-Day Delay Igniter was used in conjunction with large electric demolition charges (3, p 109-10).

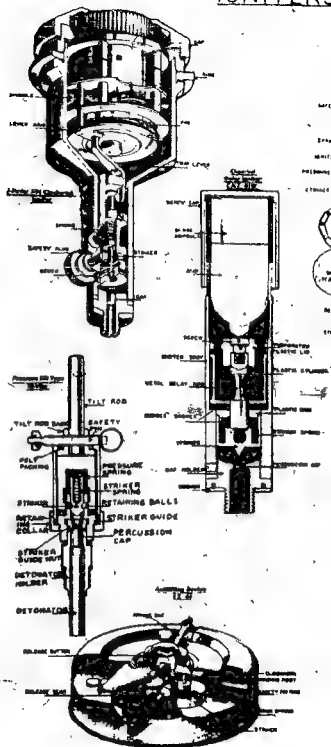
ff) 21-Day Delay Igniter was used for the same purposes as the previous igniter, but it could be set for delay ranging from 1/2 hour to 21 days. The igniter consisted of a bakelite funnel-shaped aluminum housing containing a clockwork mechanism in the upper (wide) portion and a striker assembly in the lower (narrow) portion. At the end of the predetermined time period, the lever arm on the rotating control disc bore against the trip lever, causing it to disengage the strikers. The striker, driven by a spring, exploded the percussion cap thus initiating the main HE charge (1, p 85-86 & 3, p 109-10).

Spiegelstadt

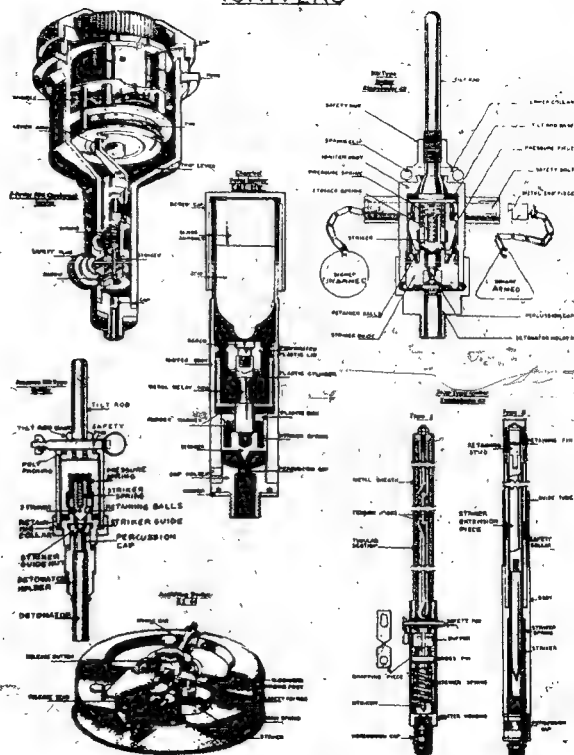
Get 97



Ge'z 58



IGNITERS



3, p 302.

K. 3 Type Igniter (Kippzunder).

a) KIZ 43 consisted of a tilt rod and a 2-4 inch extension rod connected to a cylindrical body containing the striker mechanism and a percussion cap. Lateral pressure of 25 to 25 lb exerted in any direction on the tilt rod (or 15 lb if the extension rod was used), caused the pressure piece to slide down. This allowed the retainer balls to slide outward thus releasing the striker and its spring. The impact of the striker against the percussion cap set off the explosion wave. This igniter was used in antitank and antipersonnel mines as well as in booby traps (1, p 31-376, 3, pp 313-14).

b) KIZ 45 (New Type) retained the basic principles of KIZ 43 except that it had an entirely different safety device. It is described in Ref. 1, p 31-376 and 3, pp 313-16.

1) Antitank Igniter ("Entlastungszünder"), such as E2 44 consisted of the flat cylindrical outer casing, the base plate, the clockwork and striker mechanism and the explosive filling. After winding the clockwork mechanism, the device was placed under a mine or other object and the wiring was pulled out by means of a cord or wire attached to the ring. When released, the clockwork, which ran only for 15-40 seconds, gradually forced the safety pin ring outward, thus withdrawing the safety pin. The striker was now triggered by means of the catch (seal), which in turn was held in place by the compressed spring of the release button. Release brought from the release button of the igniter allowed the striker spring to force up the rear by means of the levered stop, thus releasing the striker (2, p 163-183, 3, pp 313-16).

2) KIZ 43/1 consisted of a metallic cylindrical body and an extension composed of five tubular sections placed end to end and encased in a thin metal sheath. The extension housed five interconnecting tension hooks while the body contained the hollow striker transversely drilled above the striker pin to accommodate the cross pin to which was assembled the snapping piece. The upper end of the snapping piece engaged the lower tension hook. This igniter was designed for use in mines lying between two tracks of enemy armor or for use in which mine from the front of the functioning of the usual type igniter. The igniter opened (after removal of the safety pin) when the lateral pressure of the mine was applied to the seal and to snap at the junctions. As a result of this the tension hooks exerted a pull on the snapping piece and the striker, thus breaking the snapping piece at its weak link. This action released the spring and allowed the striker to hit the percussion cap, thus exploding the mine (2, p 163 & 3, pp 313-16).

b) KIZ 43/1 consisted of a metallic cylindrical body (housing the percussion cap, striker and spring) and a plastic tubular extension containing the plastic striker, extension, retaining stud and retaining pin. Lateral pressure on the igniter caused the tubular extension, as well as the little plastic antitank extension, to snap. This released the striker and allowed it to impact upon the percussion cap, and consequently to explode the mine. One of these mines were the same as for KIZ 43/1 (2, p 163 & 3, pp 313-16).

References:

1) Anon, Mines and Booby Traps, War Dept Field Manual FM 5-31 (1943).

2) Anon, Enemy War Materials Inventory List, Supreme Headquarters Allied Expeditionary Force (1945).

3) Anon, German Explosive Ordnance, Dept of the Army Tech Manual FM 9-1985-2 (1953).

Igniter, Bombs: According to F. Hagbaur, The Ordnance Sergeant, No. 744, p 321, the Germans employed igniter bags in all their artillery ammunition. The bag took the place of the large primers used by the U.S. Army in large and semiautomatic rounds of ammunition. The bags were either sewn to the head of the propelling charge or they were attached by means of a string. The standard substance employed in the bags during WW II was a finely grained nitrocellulose.

(See also Ignition and under Propellants)

Igniter Compositions (Zündmixture). Igniter compositions used for propellants are listed under Propellants and the igniter compositions used for Tracers are listed under Tracers.

Ignition (Zündung). Ignition of a propellant in weapons up to 50 mm was accomplished in Germany by means of a percussion cap containing black powder. Army weapons caliber 50 to 280 mm had an igniter consisting of black powder, while the naval practice in the Navy was to use powder, while the total weight of propellant. For guns larger than 280 mm an extension called Zündverschieber was used.

In addition to the primer extra igniters were sewn to both the front and rear of each section of the propelling charge.

For Flak and some Army guns the use of black powder was considered undesirable on account of its hygroscopicity and brittleness. It was reported that charges subjected to jolting consisted broken up grains which caused too rapid ignition of the propellant. Much better results were obtained on replacing black powder by a charge called Bolidung which contained Nilschilp (Nitrozellulose-Measser-Bolidung), a porous propellant prepared by leaching, with water, colloidal NC containing 8% nitrate. This propellant was also used in blank cartridges. Another replacement for straight black powder was MSP (Nitrozellulose-Schwammpulver) which contained NC 24.0, black powder 75.8 and diphenylamine 0.2%. This mixture of NC was sufficient to bind the black powder together into hard grains.

In some cases, particularly at low loading densities, where the Bolidung did not give satisfactory ignition, a Grundladung (Base charge) of special flake propellant was used. The flake was of a size intermediate between the grain charge of the tube propellant and that of the above Nilschilp.

Practically all German cannon propelling charges consisted of long tubes and were considered susceptible to ignite these at both ends. In order to ensure for the primer flash a clear passage to the front of the propelling charge, a three-walled cone shaped gas diaphragm was placed along the axis of each section of the charge. Reference: H.H.M.Pike, CIGOS Report 31-68 (1946), pp 7-8.

Ignition (Inflammation or Deflagration) Temperature (Zündtemperatur). For description of the test see: Kart-Mittel, Chemische Untersuchung der Spreng- und Zündstoffe, Braunschweig, (1944), pp 224-6 and in the general section.

The ignition temperature of some explosive and pyrotechnic compositions was determined by F. Lense, SS 27, 369-71 (1932).

(See also Flammability Test).

I.G. Wachs (IG Wax). During WW II, the I.G. Farbenindustrie developed several synthetic waxes some of which had higher melting points than natural waxes. These waxes were used for polymerizing explosives such as PETN and RDX.

Referent: A.G. Wirth, The Chemistry and Technology of Waxes, Reinhold, N.Y. (1947), p 389.

Ignitioning Compositions and Ignitioning Bombs (Leuchtstoffe und Leuchtbomben). See under Pyrotechnic Compositions and also in Stenochair, Spreng- und Schießstoffe, Zürich (1948), pp 124-9.

Incendiary Bomb. See under Bombs.

Incendiary Compositions and Incendiary Bombs. Brandstoffe oder Brandmischung und Brandbomben. According to Ref. 2, p 18) most German incendiary projectiles consisted of an Elektron (such as MgAl₂ or MgAl₂ alloy) casing filled with thermite (such as Fe oxide-70-76 and

Al 20-24%). Other fillings were white phosphorus, oil or compositions such as petroleum 87.7, polystyrene 11.9 and phosphorus 0.5% (Ref. 4, p 35). One type of projectile was prepared by filling a container with paraffin lumps of dried power pulp, followed by evaporation of oil and running in molten white phosphorus (Ref. 2, p 6). Another type, (B8) consisted of a steel outer case into which two tubes were inserted, the outer of celluloid and the inner of paper, the space between these two tubes was filled with paraffin, and the inner tube with thermite (Ref. 1, p 2).

Most incendiary bombs resembled in appearance the ordinary HE bombs. They ranged in size from 1 kg magnesium bombs (B13) to the 500 kg oil-filled-bomb (Flam 300). Several incendiary bombs were carried in a single size high explosive bomb. The 1 kg and 2 kg magnesium bombs often had a small antipersonnel charge incorporated in the bomb to discourage fire fighters. Some larger types also had a small explosive charge but this was for the purpose of wounding the incendiary mixture.

(See also Bomb: Brandbombe, Flammbombe and Sprengbrandbombe).

Only few of the German shells listed in Ref. 5 were incendiary. One of them, 50 mm HE-407 (5 cm HE-407/41 Type) was used in AA Gun, Flak 41 (377). Another was 88 mm Inc-Schrapnell (8.8 cm Gr-Schrapnell) used in AA Gun Flak 18, 36 and 47 (440).

Some German incendiaries are described by Stenochair (Ref. 2):

1) Lt. Lissawitz, RIGOS Field Report 1233 (1943), p.2.
2) W. Baumann, CIGOS Report 124-68 (1946), p. 6.
3) A. Stenochair, Spreng- und Schießstoffe, Zürich (1948), pp 124-9.

4) 197-1985-2 (1953) 5) TM 9-1985-2 (1953).

Industrial Explosives. See Commercial Explosives.

Incident-Explosion-Guidance System or Ballistic Guidance System. See under Guidance Systems.

Infant-Guidance System. See under Guidance Systems.

Infant-Red Tanning (Infant-Red Camera). Due to the fact that infrared radiation could be made invisible to the naked eye, photography, even if monochrome, coloring had been adopted, several dyes were developed by the IG Farbenwerk which were used in various types of cameras. The following types of dyes were considered to be worthy of consideration: 1) Diphenylamine 2) Diphenylamine, Carbon Black (when printed with organic dyes) and Indanthrene Olive GW 58-18.

Reference: CIGOS Report 29-218 (1945), pp 14-17.

Ignition. The same given by Dr. Valnet to hydrogen peroxide of very high concentration (such as 85%) will not be used as a fuel or as a source of stored energy. As a fuel, hydrogen peroxide superheated steam which can be used for driving other piston engines or turbines. As a source of oxygen, it was used in submarines in order to allow them to use their main engines while submerged. (See also Hydrogen Peroxide and T-Sust).

Initiating Coating. Initiated to control the burning of rocket propellant. The initiating coating was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Coating. Initiated to control the burning of rocket propellant. The initiating coating was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Coating. Initiated to control the burning of rocket propellant. The initiating coating was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Coating. Initiated to control the burning of rocket propellant. The initiating coating was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Initiating Composition. Initiated to control the burning of rocket propellant. The initiating composition was developed during WW II at the Dinsberg Fabrik, D.A.G. Its composition was polyvinyl acetate 25, lithium 30, diethylamine 10, and other organic materials. Reference: CIGOS Report 29-24 (1945), p. 5.

Igniter Compositions (Zündstoffe). Igniter compositions used for propellants are listed under Propellants and the igniter compositions used for Tracers are listed under Tracers.

Ignition (Zündung). Ignition of a propellant in weapons up to 50 mm was accomplished in Germany by means of a primer, while larger weapons required a primer combined with an igniter containing black powder. Army weapons called 50 to 700 mm had an igniter consisting of 2 g of black powder, while the navy practice in the Navy was to use 1% of black powder per total weight of propellant. For calibers larger than 280 mm an extension called Zündverstärker was used.

In addition to the primer other igniters were seen to be used, and most of each section of the propelling charge.

For Flak and some Army guns the use of black powder was considered undesirable on account of its hygroscopicity and brittleness. It was reported that charges subjected to jolting contained broken up grains which caused too rapid ignition of the propellant. Much better results were obtained on replacing black powder by a charge called Ballistol which contained NukemNH₄ (Nitrocellulose-Masovener-Nudel-pulver), a porous propellant prepared by leaching with water, cellulose NC, castor oil, and kerosene. This propellant was also used in some German rockets. Another replacement for straight black powder was HSP (Nitrocellulose-Schwamm-pulver) which contained NC 24.0, black powder 75.8 and diphenylamine 0.2%. This amount of NC was sufficient to bind the black powder together in grains.

In some cases, particularly at low loading densities, where the Ballistol did not give satisfactory ignition, the Granulit (black charge) or special flake propellant was used. The flake was of a size intermediate between the main charge of the tube propellant and that of the above NukemNH₄.

Practically all German cannon propelling charges consisted of imp-bombs and it was considered essential to ignite them at both ends. In order to ensure for the primer flake a clear passage to the front of the propelling charge, a thin-walled cordite tube of fairly large diameter was placed along the axis of the main charge. For reference H.N.M.Pike, CIGOS Report 31-68 (1946), pp. 7-8.

Ignition (Entzündung oder Zündung). Temporalzündung (Entzündungs- oder Zündungs-) Temporalzündung. For description of the test see: *Kunststoffe, Chemische Untersuchungen der Spreng- und Zündstoffe*, Reichswehr, (1944), pp. 22-6 and in the general section.

The ignition temperature of some explosive and pyrotechnic compositions was determined by F. Lenz, S. S. Z., 109-73 (1932). (See also Flammability Test).

IG Wechsels (IG Wax). During WW II, the IG Farbenindustrie developed several synthetic waxes some of which had higher melting points than natural waxes. These waxes were used for plasticizing explosives such as PETN and RDX.

Reference: A. C. Wirth, *The Chemistry and Technology of Waxes*, Reinhold, N. Y. (1947), p. 399.

Illuminating Compositions and Illuminating Bombs (Leuchtsätze und Leuchtbomben). See under Pyrotechnic Compositions and also in Suetzbecher, Spreng- und Schießstoffe, Zürich (1948), pp. 124-9.

Incendiary Bomb. See under Bombs.

Incendiary Compositions and Incendiary Bombs. Brandstoffe oder Brandstoffe und Brandbomben. According to Ref. 7, p. 18) most German incendiary projectiles consisted of an Eistrom (such as MgO) and a mixture of calcium filled with thermite such as Fe oxide-70-76 and

Al 30-240. Other fillings were using phosphorus, oil or compositions such as petroleum 87.7, polystyrene 11.8 and phosphorus 0.5% (Ref. 7, p. 2). One type of projectile was made by filling a container with peening lumps of dried paper pulp, followed by evacuation of air and running in molten white phosphorus (Ref. 7, p. 6). Another type, (B4), consisted of a metal outer case into which two tubes were inserted, the outer of celluloid and the inner of paper; the space in between these two tubes was filled with asphaltene, and the inner tube with thermite (Ref. 1, p. 7). Most incendiary bombs resembled in appearance the ordinary HE bomb. They ranged in size from 14 g magnesium bombs (B12) to the 700 kg oil-filled-bomb (Flam 500). Several incendiary bombs are listed under Bombs. The major types were usually carried in containers, whereas the larger bombs were carried in bomb racks like a single size high explosive bomb. The 1 kg and 2 kg magnesium bombs often had a small antipersonnel charge incorporated in the bomb to discourage fire fighters. Some larger types also had a small explosive charge but this was for the purpose of actuating the incendiary mixture.

(See also Bomb-Greifbombe, Flammbombe and Sprengbombe).

Only few of the German shells listed in Ref. 5 were incendiary. One of them, 50 mm HE-Inc-T (5 cm HbSpFlak 41 E type) was used in AA gun Flak 410 397. Another was 80 mm Inc-Chem-P (8 cm GrbSpFlak) used in AA gun Flak 16, 35 and 37 (p. 448).

Some German incendiaries are described by Suetzbecher (Ref. 1).

References: 1) L. Lisowski, CIGOS Final Report 1233 (1945), p. 2. 2) E. W. Bauman, CIGOS Report 32-11 (1945), pp. 6, 8, 18-19. 3) A. Suetzbecher, Spreng- und Schießstoffe, Zürich (1948), pp. 124-9. 4) TM 9-1985-2 (1953). 5) TM 9-1985-3 (1955).

Incendiary Explosives. See Commercial Explosives.

Incident Gravitation Guidance System or Ballistic Guidance System. See under Guidance Systems for Missiles.

Infar-Red Compositions. See Infar-Red Tracers.

Infar-Red Guidance System. See under Guidance Systems for Missiles.

Infar-Red Tracing (Infar-Red Compositions). Due to the fact that cloth covered objects could be readily detected by infrared photography, infar-red composites, coloring, being developed, several dyes were developed by the IG Farben which minimized or even prevented such detection. The following types of dyes were considered to be worth of consideration: Aniline Black, Diphenylamine Black, Carbon Black (synthetic with organic modern) and Indanthrene Olive GV Saffron.

Reference: CIGOS Report 25-18 (1945), pp. 14-17.

Imagin. The same given by Dr. Walter to hydrogen peroxide of very high concentration (such as 85%). Imagin can be used as a fuel or as a source of stored oxygen. As a fuel, it produces explosive steam which can be used for driving either pistons or turbines. As a source of oxygen, it was used in submarines in order to allow them to use their main engines while submerged. (See also hydrogen Peroxide and T-SO₂).

Inhibiting Coatings. In order to control the burning of rocket propellants and those for missilistic (ATO), was developed during WW II at the Döberberg Fabrik, D. A. G. The composition was polyurethane, acetone 25, diphenylamine 30, methylcellulose 5 and water 40%. Reference: CIGOS Report 10-24 (1945), p. 5.

Initiation/Initiation oder Initiationsprozess (Initiating or Priming Explosive). See Priming and Initiating Composition.

Initiators (Initiating Compositions). See Priming and Initiating Compositions.

Initiating Compositions (Initiationsexplosstoffe). See Priming and Initiating Compositions.

Initiation/Initiation (Zündkraft). The initiating property or power of primary or initiating explosives may be determined by loading as empty cap (such as the types used for 8 decaoters) with a weighed quantity of an explosive to be tested, compressing the sample and subjecting it to a load cap to one or both of the following cases: 1) Load Plane. Test of 2) Load Block Compression Test.

These tests are used for the same purpose as the American Sand Test and Nail Test, described in the general section.

Reference: A. Suetzbecher, Spreng- und Schießstoffe, Leipzig (1933), p. 134.

J (Polver). One of the sporting propellants: gunpowder, 79.5; Achromate 14.0, K bichromate 3.0; nitrate 1.5, and gelatinizer 2.5% (Brunswig, Des. zeichnung Pulver (1926) p. 134).

Jagdfeuer (Hunting or Sporting Propellant). Two kinds of propellants were used in shotgun and sporting rifles, black powder and smokeless propellant. The first successful (topical) smokeless propellant was "Schulke-Pulver". Other smokeless propellants used for sporting purposes were: Ambert, E. C. Pulver, (Polver), Saccin, and Waldox.

Reference: Brunswig, Des. zeichnung Pulver (1926), p. 134.

Jagd Feuer (Jagd Destroyer Tiger). A self-propelled mine consists of 128 mm A/S gun on PzKw VI tank under Panzer.

Jag Propellant. is briefly described in the general section. Some information on German jet units designed and manufactured by the Walter Werke, Kiel is given in CIGOS 30-15 (1945).

Jag Propellant Fuel. See under Submarine Fuel.

J-Polver-20. Clockwork top-delay (6 to 21 days) used in demolition charges [TM 9-1985-2, (1953), pp. 309-11].

Joubert. See Yonckin in the Belgian section.

Juckens Schmutzmittel. One of the gassed missiles (v) developed during WW II.

Reference: A. Dürsch, Les Armes Secretes Allemandes, Paris (1947) pp. 93-95.

Kalkammoniumsulfat (Calcium-Ammonium Nitrate) was an intimate mixture in granular form of chalk and Ammonium. It contained 20.5 to 21% N and was used as a fertilizer. Reference: R. J. Morley, CIGOS Final Rep. 289, Item 22 (1946), pp. 12-29.

Kalzipritum (Cold-priming). See Cold Extrusion in this and in the general section.

Kalstroem, Kalstroem (Cold Stretching). See Amorphization in the general section.

Kampfer (Camphor). See general section.

Kamm (K) (Cannon, Piece of Gun) Table 294 gives designations of German artillery weapons with their English equivalents.

Feldmine
 Fliegerminen
 Gebirgsmine
 Gebirgsmine
 Kampfmunition
 Kanone (Kino) (K)
 Kanone ohne Rindlauf
 leichtes Feldminen
 leichtes Kanone oder
 leichtes Geschütz
 leichtes Infanterie-
 geschütz
 Panzerabwehrkanone
 schwere Feldminen
 schwere Kanone oder
 schwere Geschütze
 schwere Infanterie-
 geschütze
 (See also under Vespene)

FK Field Gun
 Flak Antiaircraft gun
 GbH Mountain howitzer
 GbK Mountain gun
 KwK Tank gun
 KR Railroad gun
 KR Recoilless gun
 IFR Light field howitzer called by the British "gun-howitzer"
 (K)
 (IG)
 (IG) Light infantry gun
 (IG)
 Pak Antitank gun
 sFH Heavy field howitzer
 sK Heavy gun
 sKG Heavy infantry gun
 (sG)

Kanone ohne Rindlauf. See Recoilless Gun.

"Karl" Motor. See "Thor" and "Karl" Weapons.

Kartridg. See Cartridge.

KA-Salz The term assigned to RDX (Hexogen) prop by the abstraction of hexamite. An amine, nitric acid and acetic acid mixture. It is described in this section under Hexogen.

"Knacker" Target Indicating Flare. See under Flare.

Keros Explosives. Several explosive mixtures were proposed by W. Keros of W. A. S. G. One such explosive was patented in 1939 (Ref. 1). It consisted of a regulated blasting explosive plus an additional charge consisting of NG and/or nitroglycerin mixed with a large amount of alkali bicarbonate. This mixture tended to produce inert gases and to absorb heat. If desired charges containing bicarbonate could be inserted between normal charges. These explosives were suitable for use in gaseous coal mines. (See also, Bikanit and under Sheathed Explosives). Another patent granted to the same person (Ref. 2) dealt with the manufacture of moist Am nitrate explosives containing carbonaceous materials.

References:
 1) W. Keros and W. A. S. G., Brit P 493 984 (1939) C.A. 33, 2719 (1939)
 2) Ibid, Ger P 679,512 (1939); C.A. 33, 3647 (1939).

KH-Charge The designation for a compressed charge consisting of 4-8 pellets of TNT wrapped in paper glued on the inside with an adhesive glue (such as dextrin, Vinnapex, etc.). The wrapped charges were dried at 60-70° and then dipped in paraffin. They were used as bursting charges in Naval mines [See P. Rev. No 925 (1945), p. 48].

Kinetit (Kinetite). One of the oldest (1884) gelatinous explosives containing an NG gelatin. It consisted of K chlorate 37, antimony sulfide 3, nitrobenzene or nitrophenol 21 and collodion, cotton 1% Naoum, Nitroglycerin (1928), p. 553.]

King Tiger or Royal Tiger. See Königstiger, under Panther.
 -Kippender (K) (Tilt-Type Igniter). See under Igniter.

Kitchener Salt Explosives. See Kochsalzexplosstoffe.

KIAZ 40. An igniterless nose fuse used in nose rockets, such as 8.6 cm R/L (4.5) and 8.6 cm R/L (5.5). (TM 9-1985-2 (1953), p. 256.)

KMA Block. One of the substitute explosives. See under Ersatzexplosstoffe.

Knallquecksilber (Mercury Fulminate) (M F) is described in the general section under Fulminates. German methods of preparation (from mercury, nitric acid and alcohol) are given in PB Rept No 95,613 (1947), section 4. M F was used by the Germans in some painting compositions. See also A. Steinhilber, Spreng- und Schießstoffe, Zürich (1949), pp. 95-96.

Knallpulver (Silver Fulminate). See general section under Fulminates and Steinhilber's book (1949) p. 96.

Knallschloß (Detonating Fuse). See general section under Fuses.

Knetmaschine (Kneading Machine). An apparatus used for mixing solid ingredients in the presence of liquids. Several types were used in Germany such as the Coleman type (Glasbleichmaschine) (Ref. 2, pp. 105, 106 and Ref. 3, p. 237), Vetter-Pfeifferer-Maschine and Knetmaschine (Ref. 1, p. 75 and Ref. 3, p. 227) and others.

References:
 1) E. de B. Barnett, Explosives, Van Nostrand, N.Y. (1919)
 2) P. Steinhilber, Schieß- und Sprengstoffe, Stuttgart, Dueden (1927)
 3) A. Steinhilber, Schieß- und Sprengstoffe, Barth, Leipzig, (1935).

Kneisender 43 (Snop Type Igniter). See under Igniter.

Kochsalzexplosstoffe (Kitchen Salt Explosives). Substitute explosive mixtures containing large amounts of Na chloride, which were used during WW II. Some of these mixtures are described under Ersatzexplosstoffe.

Kohlen-Carbonit
 Kohlen-Korant III
 Kohlen-Salz
 See under Kohlenexplosstoffe.

Kohlenexplosstoffe (Coal Explosives). This was a group of explosives permitted for use in coal mines.

Kohlen-Carbonit NG 25, K nitrate 34, Bp nitrate 1, flour 38.5, spent tan meal 1 and soda ash 0.5%; heat of explosion 306 kcal/kg; temp of explosion 1561°C, velocity of detonation 3160 m/sec, density 1.16 and Trauzl test value 235 cc (Ref. 2, p. 401 and Marshall, 1942, p. 492).

Kohlen-Korant III NG 4, K chlorate 68, Na chlorate 14, paraffin 8, nitrophenol 3 and wood meal 1%; oxygen balance -12% and Trauzl test value 195 cc (Ref. 3).
 Kohlen-Salz NG (gelatinized) 12.5, meal 2.5, nitrocompounds 7.0, Am nitrate 47.0 and alkali chloride 37.0%; oxygen balance -2.6% and Trauzl test value 260 cc (Ref. 2, p. 441).

Kohlen-Wassalf II NG 4.0, Am nitrate 63.0, K nitrate 7.0, Bp nitrate 2.0, meal 2.0 and TNT 2.0%; oxygen balance 216.4% and Trauzl test value 230 cc (Ref. 2, p. 415).

Kohlen-Wassalf IV NG 3.2, Am nitrate 75.0, K nitrate 2.8, alkali chloride 15.0, meal 1.0, and DNT 5.0%; oxygen balance -8.8% and Trauzl test value 200 cc (Ref. 2, p. 415).

Kohlen-Wassalf V NG 4.0, Am nitrate 85.0, K nitrate 8.0, Bp nitrate 2.0, potato meal 1.5 and Montana was 1.5%; oxygen balance +13.5% and Trauzl test value 230 cc (Ref. 2, p. 415).

References:
 1) P. Steinhilber, Schieß- und Sprengstoffe, Dueden (1927), p. 147
 2) P. Steinhilber, Nitroglycerin, etc., Baltimore (1928), pp. 435 & 441.

Kohlen-Wassalf. See under Kohlenexplosstoffe.

Kohlen-Tin explosive of the carbamate type, such as NG 23, K nitrate 75, Bp nitrate 5, wood meal 34, and starch 10%. There was also a Super-Kohlen, an explosive used in England [Marshall 1 (1917), p. 375].

Kollie (Kollite). A smokeless propellant patented in 1900 by H. Kollie of Berlin consisted of mixtures of nitrocellulose, nitro, oil cakes, residues of factories manufacturing organic products such as starch, sugar, beer, alcohol, etc., with saltpetre previously saturated with nitrobenzene.

References: J. Daniel, Dictionnaire, Paris (1902), p. 394.

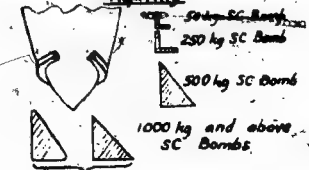
Kontinuierliche Verfahren (Continuous Methods) of manufacture of explosives such as those of Schmid, Meissner and Blum were used in several German plants.

Some of these methods are briefly described under Nitroglycerin, Korant and Trinitrophenol, as well as in the Belgian, Dutch, French, Swedish and Swiss sections.

References:
 1) A. Steinhilber, Schieß- und Sprengstoffe, Barth, Leipzig (1935), pp. 174 & 313
 2) A. Steinhilber, Spreng- und Schießstoffe, Barth, Zürich (1949), pp. 60 & 97
 3) A. Steinhilber, Potentia Explosives, Gill, Bureau (1952).

Koppling (Head Ring). When it was desired to avoid excessive protrusion against head target and to prevent ricochet against the target, rings were attached over the nose of bombs such as SC (HE cylindrical, general purpose) or SC 200 (A/P cylindrical, thick walled). (See also Anti-Aircraft Flares).

References: TM 9-1985-2 (1953), p. 3.



Korant V. On during WW I. In 19, nitrophenol and Sprengstoffe. Note: According to Korant was

K Pulver. St

Kraftstoff (KZ) of power (strength) Lead Block TNT error is due which is observed in gelatin, fire error, Neubauer expansion process the weight of firing several obtain values curve giving up expansion to be determined is called Kraftstoff. Table 25

Substance
Blasting gelatin
NG
NCI (35)
P A
TNT
DNS

Note: It may KZ values be as determined potential explosive higher.

References:
 1) R. Neubauer,
 2) A. Steinhilber,
 3) A. Steinhilber

"Kronit". A solid blasting Reference: TM

Kropfdruck (Cr

Kropfdruck (Cr) of power (strength) in "Flak" is fixed maximum 31-62, p. 53.

Kropfdruck (Cr) Double base Nitro Pulver (KOP) 31-62

King Tiger or Royal Tiger. See Koolgastig, under Plastic.

Kippzunder (J). (Tri-Type Igniter). See under Igniter.

Kitchen Salt Explosives. See Kochsalzexplosstoffe.

KIAZ 40. An impact-firing nose fuse used in some rockets, such as 8.6 cm R/L(4.5) and 8.6 cm R/L(5.5). (TM 9-1983-2 (1953), p 716).

KMA Block. One of the substitute explosives. See under Ersatzexplosstoffe.

Knallquecksilber (Mercury Fulminate) (M F) is described in the general section under Fulminates. General methods of preparation (from mercury, nitric acid and alcohol) are given in PB Rept No 93.63 (1947), section 4. M F was used by the Germans in some priming compositions. See also A. Stuebner, Spreng- und Schietstoffe, Zürich (1948), pp 95-96.

Knallsilber (Silver Fulminate). See general section under Fulminates and Stuebner's book (1948) p 96.

Knallzunder (Detonating Fuse). See general section under Fuses.

Knetmaschine (Kneading Machine). An apparatus used for mixing solid ingredients in the presence of liquids. Several types were used in Germany such as the Columnar Type (Staubenmischmaschine) (Ref 2, pp 105, 106 and Ref 3, p 373). Werner-Pfeiderer Misch- und Knetmaschinen (Ref. 1, p 75 and Ref 3, p 227) and others.

References:
1) E. de B. Barnett, Explosives, Van Nostrand, N.Y. (1919)
2) P. Noxon, Schiess- und Sprengstoffe, Sorikopi, Dresden (1927)
3) Stuebner, Schiess- und Sprengstoffe, Bern, Leipzig, (1933).

Knetzunder (J) (Soap Type Igniter). See under Igniter.

Kochsalzexplosstoffe (Kitchen Salt Explosives). Substances explosive mixtures containing large amounts of Na chloride, which were used during WW II. Some of these mixtures are described under Ersatzexplosstoffe.

Kohlen-Carbonit
Kohlen-Kornit H. See under Kohlenexplosstoffe.
Kohlen-Salt
Kohlen-Salt

Kohlenexplosstoffe (Coal Explosives).

This was a group of explosives permitted for use in coal mines.

Kohlen-Carbonit. NG 25, K nitrate 34, Ba nitrate 1, flour 38.5, spent gun metal 1 and rock ash 0.5%; temp of explosion 306 kcal/kg; temp of explosion 1961°C; velocity of detonation 3160 m/sec; density 1.16; and Trauzl test value 235 cc (Ref 2, p 401 and Marshall, 2, p 492).

Kohlen-Kornit. 81.0% Na K chlorate 68, Na chloride 14, paraffin 8, nitrophenol 5 and wood meal 1%; oxygen balance -12% and Trauzl test value 195 cc (Ref 1).
Kohlen-Salt. NG (gelatinized) 12.5, meal 2.5, nitro-compounds 7.0, Am nitrate 41.0 and alkali chloride 37.0%; oxygen balance -2.6% and Trauzl test value 260 cc (Ref 2, p 441).

Kohlen-Wasserdampf 1. NG 4.0, Am nitrate 83.0, K nitrate 7.0, Ba nitrate 2.0, meal 2.0 and TNT 2.0%; oxygen balance 63.6% and Trauzl test value 230 cc (Ref 2, p 435).

Kohlen-Wasserdampf JV. NG 3.2, Am nitrate 73.0, K nitrate 2.0, alkali chloride 15.0, meal 1.0, and DNT 5.0%; oxygen balance 18.8% and Trauzl test value 300 cc (Ref 2, p 435).

Kohlen-Wasserdampf V. NG 4.0, Am nitrate 83.0, K nitrate 8.0, Ba nitrate 2.0, potato meal 1.5 and Monax 1.5%; oxygen balance +13.5% and Trauzl test value 230 cc (Ref 2, p 435).

References:
1) P. Noxon, Schiess- und Sprengstoffe, Dresden (1927), p 147
2) P. Noxon, Nitroglycerin, etc., Baltimore (1928), pp 415 & 441.

Kohlen-Wasserdampf. See under Kohlenexplosstoffe.

Kornit. An explosive of the carbonate type, such as: NG 25, K nitrate 35, Ba nitrate 5, wood meal 34, and starch 10%. There was also a Super-Kornit, an explosive used in England (Marshall 1 (1917), p 375).

Kornit (Kornit). A smokeless propellant patented in 1890 by H. Kohn of Bonn, which consisted of mixtures of nitrocellulose, nitro, oil colors, residues of faceted manufacturing organic products such as starch, sugar, beer, alcohol, etc., with talcaper previously saturated with nitrobenzene.

Reference: J. Daniel, Dictionnaire, Paris (1902), p 394.

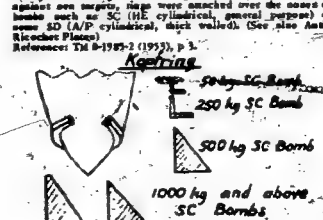
Kontinuierliche Verfahren (Continuous Methods) of manufacture of explosives such as those of Schmid Meissner and Bissel were used in several German plants.

Some of these methods are briefly described under Nitroglycerin, Picramit and Trinitrotoleol, as well as in the Belgians, Dutch, French, Swedish and Swiss sections.

References:
1) A. Stuebner, Schiess- und Sprengstoffe, Bern, Leipzig (1933), pp 174 & 335
2) A. Stuebner, Spreng- und Schietstoffe, Bacher, Zürich (1948), pp 60 & 87
3) A. Stuebner, Poisons Explosives, Gill, Bombay (1952).

Koppling (Head Ring). When it was desired to avoid excessive penetration against lead targets and to prevent ricochet against one target, rings were attached over the nose of some work or SC (i.e. cylindrical, general purpose) or some SD (A/P cylindrical, stick value). (See also Anti-Ricochet Plans).

Reference: TM 9-1983-2 (1953), p 3.



Kornit. V. One of the permissible explosives developed during WW I: NG 4, K chlorate 65, Na chloride 15, nitrophenol 5, and wood meal 2%. (Noxon, Schiess- und Sprengstoffe, Dresden (1927), p 147).
Note: According to Marshall, v 3 (1932), p 112, the name Kornit was given in 1931 to Chloratit 1.

K Pulver. Same as G Pulver.

Kraftzahl (KZ) (Strength Number). In the usual determination of power (strength) of an explosive by the standard Trauzl Lead Block Test, (see general section) one of the principal errors is due to weakening of the walls of the cavity, which is observed with powerful explosives such as blasting gelatin, P.A., TNT and NG. In order to eliminate this error, Neuber proposed that, instead of measuring the expansion produced by a standard weight of explosive, the weight of explosive required to produce a standard expansion of 300 cc be determined. This may be done by firing several charges of different weights in order to obtain values below 300 cc and above it. After drawing a curve giving the relationship expansion vs weight of sample, the expansion in cc corresponding to a 10 g sample can be determined by interpolation. This calculated expansion is called Kraftzahl (strength number).

Table 25b lists KZs for some explosives

Substance	Trauzl Test: Volume observed by various investigators using a 10 g sample	KZ calculated by Neuber for a 10 g sample
Blasting gelatin	520 to 610cc	55-60
ENG	315 to 600	540
NC(13%)	325 to 420	400
P.A.	300 to 360	380
TNT	225 to 300	350
DMB	250	311

Note: It may be concluded from the above values that the KZ values for highly powerful explosives are lower than are determined by the standard Trauzl test, while for less powerful explosives (such as TNT, or DMB) the KZ is higher.

References:
1) R. Neuber, S.S. 23, 54 (1928)
2) A. Stuebner, Explosives, v 3 (1932), p 143
3) A. Stuebner, Spreng- und Schietstoffe (1948), p 113.

"Kornit". An accurate proximity fuse intended for some guided missiles as, for instance, Rocket X-4.

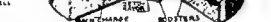
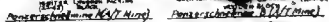
Reference: TM 9-1983-2 (1953), p 216.

Kornit (Crown Bomb). See Distance Fuse.

Kornit (Kornit) Pulver. Double-base DEGN-NC propellant with a caloric value of 710-730 kcal/kg, used in flash rockets in lieu of K sulfate used in G Pulver (CDS 31-62, p 53).

Kornit (Pulver) also Nitrat aber mit Dinitrotoleol (KOD). Double base DEGN-NC propellant similar to Kornit Nitrat Pulver except that K nitrate was replaced by DNT (CDS 31-62, p 53).

Get 10%



6. 106.

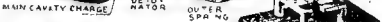


Fig. 3. Details of the hoist mechanism.

Marine Explosives of WW I and WW II. Under this title, A. Stettbacher (Switzerland), 9, 33-45 (1943), describes the explosives used by the Germans for filling torpedoes, sea mines, depth charges etc:

- Explosive of WW I: TNT 60, HNDPHA (hexanitrodiphenylamine) 40%
- Explosive of WW II: TNT 61.8, HNDPHA 23 and Al powder 15.1%

The second mixture was much more effective than the first one.

Marine-Geschütz Pulver. Black powder used as a bursting charge in phosphor bombs, such as BL 7.5-20 bombs. The composition of the powder was: K nitrate 75, sulfur 9 and beach charcoal 16%. The granulosity was 0.68 to 1.3 mm and the moisture content 1.3%.

Reference: TM 9-1985-2 (1953), p. 82.

Mark 50 Kerosene (Candle) Flame Bomb. See under Pyrotechnic Anti-Peashooter Devices.

Marker. (Azeigier). A pyrotechnic device used to mark a position. Most of the German markers consisted of cylindrical cardboard containers filled with a colored flare composition which was ignited by an impact type fuse. Some markers merely contained a brightly colored powder, which was dropped into the sea from low altitudes to mark positions. Others were incendiary.

The following devices described in TM 9-1985-2 (1953), could be classified as markers:

- NC 50 NC 1/35 SE Smoke Marker Bomb resembled an ordinary HE bomb. It consisted of an aluminum outer casing (tepy except for metal ribs and braces), tail cone, nose and central cylinder which extended from the nose and extended aft to the forward part of the tail where it was terminated by a flare housing composed of 1. Waterproofing at the tail was provided by a rubber seal. The central cylinder contained the smoke producing agent. Four fins and a plate (called drogue) were attached to the tail end, impact of the bomb on water caused the drogue, together with the fuse release rod, to be wrench off. This action fired the fuse and ignited the smoke mixture. At this time the bomb would be floating on the surface. Eventually the heat from the burning smoke composition destroyed the rubber seal and the smoke was vented to the outside, thus indicating the position of the marker. (pp 59-60)
- Mark 5 Flare, Types 3 and 4. Floating devices which could serve as markers or for signaling purposes. (See under Flare and in TM 9-1985-2, pp 73-74)
- Target Indicator (Red) consisted of an aluminum cylindrical casing housing a flare composition enclosed in a cardboard cylinder. The suspension flare at the tail held an eye to take the parachute shackle, and a pull igniter which was connected by a 4 1/2 inch length of safety fuse to a burning cone black powder. This served both to set off the igniter pellet in the top of the candle, and to eject the latter from the casing when it fell freely to earth and acted as a goosene marker. The pull igniter was attached to the loop of the shroud lines by a cord and the opening of the parachute gave the necessary pull for operating the igniter. There were (for some unknown purpose) two small fins at the nose end of the container. (pp 64-65)
- See Marker Bomb consisted of sheet steel, bomb shaped container, supported internally by a series of annular strengthening bulkheads. The tail end of the bomb was provided with four stabilizing fins and a drogue housing a lamp unit covered with a lucite dome. A battery of six dry cells was housed in the center of the bomb. At the moment of the release of the bomb from the aircraft, the inertia bolt was positioned between the plates of the spring switch in such a manner that one side of the circuit between the lamp and the battery was broken. On impact of the bomb, the inertia bolt was forced out of position and the circuit between the lamp and battery was completed. As the battery was filled only a portion of the bomb body and as all joints were made tight by rubber washers, the marker floated on the surface of water. It was assumed that the marker provided a recognition or bearing point for

aircraft. (pp 83-84)

5) Sea Marker Lux 82 40 SC was constructed of sheet steel in two parts (nose and tail) loosely joined together about 1/2 inch apart from the nose. The interior view and a brief description are given on p 87 of TM 9-1985-2 (1953).

6) Mark 3 Gun (Single Unit Ground Marker, Green) consisted of a sheet steel cylinder enclosing a cardboard container with the pyrotechnic composition, a fuse with pin (filled with black powder) in an aiming spindle and an aiming vane, which was loosely fitted within the housing. On release of the marker from the aircraft, the current of air rushed through the vane holes in the aiming vane, thus ejecting it from the housing. By reason of its shape, the aiming vane rotated as the missile fell. The black powder in the spindle became ignited. The resulting flash ignited the pyrotechnic composition and at the same time a slight explosion took place which ejected the cover cap, late and aiming vane housing. The pyrotechnic filling burned for about 3 1/2 minutes.

7) Lux 3 Flame Float. A bomb-like device constructed of sheet steel and provided with four fins. When released over water the device went undriven the surface thus allowing the water to enter the ports and to pass down the central tube into the calcium phosphide chamber. The resulting reaction produced phosphine gas which passed up the outer tube through numerous vertical holes in the burner where it ignited spontaneously to form a pilot jet. At the same time, water entered through the channels in the nose and passed through a perforated tube into the calcium carbide compartment. The acetylene evolved passed through the perforated diaphragm into one compression chamber and thence to the burner where it was ignited by the pilot jet. (pp 91-92)

8) Lux 5 Flame Float (Types 1, 2 and 3) was cylindrical in shape and contained, as in the previous device, Ca phosphide and Ca carbide. (pp 92-93).

Merapelle or Mers Priming Drogs. Low tension fuseheads intended for ordinary instantaneous detonators. They were manufactured by dipping the tip of the electric bridge wire into the following liquids:

- 1st dip composition consisted of 100 g of dry Pb picrate suspended in 50 ml of a 2% solution of NC in amyl or butyl acetate. After the drop on the tip became dry it was dipped into
- 2nd dip composition consisting of Pb picrate 40 g, K perchlorate 35 g and alderwood charcoal 25 g, suspended in 50 ml of a 2% solution of NC in amyl or butyl acetate
- 3rd dip composition contained K perchlorate 85.7 and alderwood charcoal 14.3 g, suspended in about 50 ml of a 3% solution of NC in amyl or butyl acetate
- 4th dip composition was a lacquer consisting of a 15% solution of NC in 75/25 butyl acetate which was added (20% of the dry weight of NC) Sipiñol AOM which is the methylcyclohexyl ester of adipic acid, and 17 g of Sudan Brown for each 10 l of liquid.

Note: A few materials to be used in tropical countries, the 4th dip consisted of Al powder (200 g per liter of lacquer) which was supported to protect the fusehead against static electricity.

B) Merapelle contained the property of not igniting fireproof, which was a great advantage.

C) The soldering of the bridge wire to lead-in wires, the preparation of dry head dips, the preparation of NC varnishes and the process of dipping the fuseheads are described under Fusehead Manufacture.

References:

- 1) BLOS Final Rept No 853, Item 21(946), p 35/36
- 2) PB Rept No 35,613 (1947), Section D.

Messungsmessgerät (Machine Gun) See under Weapons.

Messing (Messel). A heavy metal designed by Potach (See Experimental Tanks, under Panzer)

Messing (Messel). One of the WW I straight dynamites: NC 60.0, Nitro glycol pulp 10.0, nitroed ivory nut meal (coconut) 10.0 and Na nitrate 20.0% [P. Neosim, Nitroglycerin, Baltimore (1928), p 284.]

Messingpulver (Meal Powder). Finely pulverized black powder used in pyrotechnic compositions. Its preparation is described by A. Stettbacher, Schaeus- und Sprengstoffe, Leipzig (1933), p 103 (See also Meal Powder, in the general section).

Messing. A jelly originally prepared by Sprengstoff A.G. Carbonsäure, Schickler, by adding glycerin with an aqueous solution of glue. It was incorporated in some dynamites in order to increase their plasticity. Some glycerin-glue mixtures contained dextrin (See also Glycerin-Carbonsäure and Safety Jelly Dynamite).

Reference: P. Neosim, Nitroglycerin, Baltimore (1928), p 406.

Messingboxen (Message Container or Message Tube). A device for dropping messages. Two types of containers used for this purpose are described in TM 9-1985-2 (1953), pp 120-121.

a) See Message Tube consisted of an aluminum cylinder in which the upper compartment contained a smoke composition, whereas the lower (striking) compartment carried a message. On dropping the missile from the plane, the friction igniter was pulled and the resulting flash ignited the dye flare, which in turn ignited the bottom part of the smoke composition, when persons for whom the message was intended, saw the smoke, they approached the missile and removed the message container by opening the cap (at the rear of the tube) and pulling the chain (p 120)

b) Land Message Tube was also cylindrical in shape and consisted of two compartments. The smoke composition in the upper compartment was ignited by means of a few strands of quickmatch which extended down the side of the smoke container and met several pieces of quickmatch below the striking igniter. The strands were ignited when the friction igniter was pulled on dropping the missile from a plane. The message was then picked up by pulling the nut and rearing the cap cover. (p 121)

Messing Pulver. See Kalkpulver.

Messing, Flamm Granade (Nachrichtens Piktogrammatische See under Flamm Granades.

Message Tube. See Meldeboxen.

Messel (Measuring Egg). A device designed by the Knapp plant for measuring the pressure developed in pure, the amount to which a copper cylinder was compressed by the gases of combustion of a propellant served as a measure of the maximum pressure developed in chamber. For more information on this subject, see H. Braunig, Das rauchlose Pulver, Berlin (1926), p 412.

Messingpulver. Trade name for m-Toluenesulfonamide, H₂C₆H₄SO₂NH₂ · white crystals, m.p 107°. Its solution in some organic media was claimed to be a good gelatinizer. (See NC)

Messingpulver. Trade name for m-Toluenesulfonamide, H₂C₆H₄SO₂NH₂ · white crystals, m.p 107°. Its solution in some organic media was claimed to be a good gelatinizer. (See NC)

Messingpulver. Trade name for m-Toluenesulfonamide, H₂C₆H₄SO₂NH₂ · white crystals, m.p 107°. Its solution in some organic media was claimed to be a good gelatinizer. (See NC)

Messingpulver. Trade name for m-Toluenesulfonamide, H₂C₆H₄SO₂NH₂ · white crystals, m.p 107°. Its solution in some organic media was claimed to be a good gelatinizer. (See NC)



See Messing

Messingpulver

Messingpulver

Messingpulver

Messingpulver

Messingpulver

Messingpulver

Messingpulver

Messingpulver

Messingpulver

Messingpulver

auctral (pp 85-6)

3) See Marker LUX LZ 50 SC was constructed of sheet steel in two parts (nose and tail) loosely joined together about 1/2 inch distance from the nose. Its external view and a brief description are given on p 87 of TM 919852 (1953).

6) Mark 3 Gun (Single Unit Ground Marker, Green) consisted of a sheet steel cylinder enclosing a cardboard container with the pyrotechnic composition, a fuse with gaine (filled with black powder), an arming spindle and an arming vane, which was loosely fitted within the housing. On release of the marker from the aircraft, the current of air rushed through the vent holes in the arming vane, thus ejecting it from the housing. By reason of its shape, the arming vane rotated as the missile was falling. This rotation unscrewed the arming spindle of the fuse thus permitting its clockwork mechanism to function. At the expiration of predetermined delay, the black powder in the gaine became ignited. The resulting flash ignited the pyrotechnic composition and at the same time a slight explosion took place which ejected the cover cap, fuse and arming vane housing. The pyrotechnic filling burned for about 1/2 minutes.

2) Lux 5 Flame Float. A bomb-like device constructed of sheet steel and provided with four fins. When released over water the device went under the surface thus allowing the water to enter the orifices and to pass down the inlet tube into the calcium phosphide chamber. The resulting reaction produced phosphine gas which passed up the outlet tube through the nonreturn valve to the burner where it ignited spontaneously to form pilot jet. At the same time, water entered through the channels in the nose and passed through a perforated tube into the calcium carbide compartment. The acetylene evolved passed through the perforated diaphragm into one compression chamber and thence to the burner where it was ignited by the pilot jet (pp 91-2).

8) Lux 5 Flame Float (Types 1, 2 and 3) was cylindrical in shape and contained, as in the previous device, Ca phosphide and Ca carbide (pp 92-3).

Marsalle or Mars Priming Drops. Low tension fuseheads intended for ordinary instantaneous detonators. They were manufactured by dipping the tip of the electric bridge wire into the following liquids:

a) 1st dip composition consisted of 100 g of dry Pb picrate suspended in 50 ml of a 2% solution of NC in amyl or butyl acetate. After the drop on the tip became dry it was dipped into

b) 2nd dip composition consisting of Pb picrate 40 g, K perchlorate 35 g and alderwood charcoal 25 g, suspended in 50 ml of a 2% solution of NC in amyl or butyl acetate.

c) 3rd dip composition contained K perchlorate 85.7 and alderwood charcoal 14.3 g, suspended in about 50 ml of a 3% solution of NC in amyl or butyl acetate.

d) 4th dip composition was a lacquer consisting of a 15% solution of NC in 75/25 butyl acetate ethanol to which was added (20% of the dry weight of NC) Sudan AOM (which is the methylcyclohexyl ester of adipic acid) and 1% of Sudan Brown for each 10 l of liquid.

Notes: A) For materials to be used in tropical countries, the 4th dip contained Al powder (200 g per liter) of lacquer, which was supposed to protect the fusehead against attack of the insects.

b) Methylpicric possessed the property of not igniting freudend, which was a great advantage.

c) The soldering of the bridge, were to lead-in wires, the preparation of dry ingredients for fusehead tips, the preparation of NC yarn and the process of dipping the fuseheads are described under Fusehead Manufacture.

References:

- 1) B I O S Final Rept No 833, Item 2(1946), p A3/36
- 2) PB Rept No 95,613 (1947), Section D.

Maschinengewehr (Machine Gun) See under Weapons.

"Mouse" (Mouse). A heavy cap designed by Porsche (See Experimental Tanks, under Panzer)

Megaton (Megaton). One of the 1/4 lb straight dynamites. NC (60), brisant wood pulp 10.0, nitrocellulose 10.0 (control) 10.0 and Ne minute 20.0% (P.Naoum Nitroglycerin, Baltimore (1928), p 284).

Mehlpulver (Meal Powder). A finely pulverized black powder used in pyrotechnic compositions. Its preparation is described by A. Seestrich-Schwarz and Sprengstoffe, Leipzig (1933), p 103 (See also Meal Powder in the general section).

Melma. A jelly originally prepared by Sprengstoffe A-G Claiborn, Schleibach, by adding glycerin with an aqueous solution of glue. It was incorporated in some dynamites in order to increase their plasticity. Some glycerin-glue mixtures contained dextrin (See also Gelatine-Carbonit and Safety Jelly Dynamite).

Reference: P.Naoum, Nitroglycerin, Baltimore (1928), p 406.

Meldebüchse (Message Container or Message Tube). A device for dropping messages. Two types of containers used for this purpose are described in TM 919852 (1953), pp 120-1:

a) See Message Tube consisted of an aluminum cylinder in which the upper compartment contained a smoke composition, whereas the lower (straight) compartment carried a message. On dropping the missile from a plane, the friction igniter was pulled and the resulting flash ignited the delay fuse, which in turn ignited the bottom part of the smoke container. When persons for whom the message was intended, saw the smoke, they approached the missile and removed the message container by opening the cap (at the rear of the tube) and pulling the chain (p 120).

b) Lead Message Tube was also cylindrical in shape and consisted of two compartments. The smoke composition in the upper compartment was ignited by means of four strands of quickmatch which extended down the side of the smoke container and met several pieces of quickmatch below the smoke container. The strands were ignited when the friction igniter was pulled on dropping the missile from a plane. The message was withdrawn by unlocking the nut and removing the cover (p 121).

Mercure Fulminate. See Knallquecksilber.

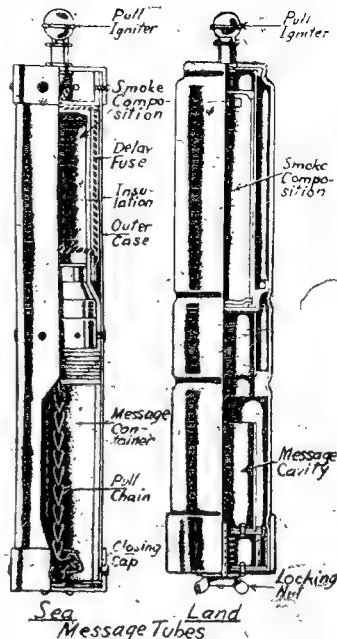
Messings Pistol Granade (Nachrichten Pistolengranate). See under Pistol Granades.

Messung Tube. See Meldebüchse

Messner (Measuring Egg). A device designed at the Knippl plant for measuring the pressure developed in gases. The extent to which a copper cylinder was compressed by the gases of combustion of a propellant served as a measure of the maximum pressure developed in chamber. For more information on this subject, see H. Braunwig, Das rauchlose Pulver, Berlin (1926), p 412.

Methocellulose. Trade name for α -D-glucosulfonamide, $H_2C_2H_4SO_3NH_2$; white crystals, m.p. 107°. Its solution in some organic media was claimed to be a good gelatinizer for NC.

Reference: Kaut-Metz, Chemische Untersuchungen, Braunschweig (1940), p 162.



Methylamine. Its preparation and properties are given in the general section. According to Dr. H. Walter, methylamine was never used in Germany per se but in the form of its nitrate, called Mas-sals (qv).

Methylamine Nitrate See Mas-Sals.



Methylnitropropanediol Dinitrates. described in the general section, was examined in Germany during WW II as a possible substitute for NG in propellants. It was found to be fairly stable but not a very good gelatinizer for NC.

Reference: PB Rept 925 (1941), p 15.



SIÈGE FICHE 273

3414-4

PICATINNY ARSENAL
TECHNICAL REPORT NO. 2510

DICTIONARY OF EXPLOSIVES, AMMUNITION AND WEAPONS

(GERMAN SECTION)

BASIL T. FEDOROFF

HENRY A. AARONSON

GEORGE D. CLIFT

EARL F. REESE

DOVER, NEW JERSEY

1958

Methylal. A mixture of aluminum dichloromethyl, $AlCl_2 \cdot CH_3$, and aluminum chloromethyl, $AlClH_2$, proposed as a substitute for phosphorus in incendiary compositions. The mixture was prepared by passing methyl chloride vapor through copper-free aluminum turnings. Reference: R.E. Richardson et al., CIOB 25-18 (1945), p. 45.

Methyltrinitrate (Methyl Trinitrate). (Methyltrinitrocellulose Trinitrate or Pentaerythritol Trinitrate), $H_2C(O_2NO)_3$, described in the general section, was developed in Italy before WW II by Bombieri Parodi-Delfino and adopted later by the Germans.

The following method of preparation was used at the Krimmel Fabrik of D.A.G.:

a) 50 kg of finely pulverized and sieved Methyl was slowly fed with stirring by means of a worm screw, into a stainless steel nitrate containing 175 kg of nitric acid, (65% HNO_3 and 35% H_2SO_4) maintained at 40°. Formation of lumps had to be avoided because this could lead to overheating and decomposition of methyl and acid.

b) After 20 minutes of nitration, 15 minutes were allowed for separation of the oil from the acid.
c) The separated oil was washed, first with water, then with soda ash solution and finally with water. The temperature during all the washings was maintained at 40° because at a low temp the mixture was too viscous. The soda ash wash lasted for 20 minutes. The yield was 300 parts Methyl per 100 p of the acid.
d) The washed oil was taken to a storage tank from which it was withdrawn when needed for the preparation of "Rohpulvermischung" (Rawpaste) (q.v.).

German technical Methyl was a heavy oil, practically insoluble in water, with the following properties: $n_{D,20}^{20} = 1.465$, $d_{4,20}^{20} = 1.465$ at 20°, stability by Abel test at 82° 20 min, decomposition temperature ca. 182°, impact sensitivity with a 2 kg hammer 4 cm; caloric value 1270 kcal/kg (water in liquid phase), volatility less than NG.

It was used in some amoniacal propellants as a explosive plasticizer for NG in the form of NG.

Reference: P.B. Rept 923 (1945), pp 15 & 16.

Mittelzink (Miedziankin). A type of chlorine explosives manufactured in Germany and Poland before WW II: a) K or Na chlorate 80-9% and liquid, hydrous acid (with flash point not below 10°C) 12-9% (Ref. by b) c) chlorate 50 and petroleum 10%. The first mixture belonged to the group of Chlorates 3.

References:

- 1) P. Noon, Shieser und Sprengstoffe (1927), p. 131
- 2) A. Szentbachi, Spreng- und Schießstoffe (1940), p. 91.

Mikrowärmeröhre beim Sprengen (Microdelay in Blasting) is described by Z. Peitner, Explosivstoffe 1954, Heft 5/6, pp 68-70.

Mine, Land. See Landmine.

Minehund/Mine Dog, called by the Allies "Doodley" or "Goldie", was a miniature remote-control tank operated by remote control through a 750 ft 3 strand cable which was wound from a drum on the tankette. Separate electric motors, each powered by its own storage battery, drove the tracks of the tank at 4 mph. Steering was done by breaking the tracks. The tank consisted about 250 kg. HE demolition charge which the remote-control operator was supposed to touch off after stopping the vehicle at its target.

These mobile mines were not very effective because they could not move in reverse. On account of their low speed and thin armor, they were easily destroyed by the Allies' artillery.

Reference: Anon, Field Artillery Journal 34 505. (1944).

Minotaur Torpedoes. See Explosive Powered Vortices.

Mining Effect. See Earth-Displacement Test.

Mining Explosives. See Commercial Explosives.

Mipolan and Mipolan Sealing Plug. Mipolans are plastic compositions developed in Germany during WW II and used in the repair of seals for some delay detonators. Previous to WW II fired seals were used. The Mipolan sealing plugs were made in three types:

- a) Long grayish-green plug with a single hole
- b) Short grayish-green plug with two holes. The Mipolan was composed of polyvinyl chloride 50, tricresyl phosphate 30 and talcum 20%.
- c) Short red-dish plug with two holes. The Mipolan was composed of polyvinyl chloride 51, Special Mixture 51, and talcum 18%.

Note: The Special Mixture consisted of 2 parts tricresyl phosphate 2 per Palatinol HC and 2 per Palatinol K. The composition of Palatinol HC was not given, and the Palatinol K was butyleneglycolphosphate.

Mipolan was also used for covering the lead-in wires of electric detonators. The thickness of coating for 60 mm wires was only 0.25 mm on detonators not intended for underwater operations and 0.35 mm on those intended for such operations.

References:

- 1) W. Kramlich, Kunststoffe in technischen Kerosinmaschinen, Lehmann, Berlin (1943), p. 23
- 2) B.I.O.S. Rept (Final), No 833, Item No 2, London (1946) on PB Rept No 63, 677 (1946)
- 3) P.B. Rept No 95,613 (1947), Sections II, I, and J.

Note: According to M.F. Fogler et al., CIOB Rept 21-5 (1945), p. 5 there were three types of Mipolan: a) Plasticized polyvinyl chloride b) Copolymers of polyvinyl chloride and acrylic ester and c) Polyvinyl chloride and maleic ester.

Mischmetall (Mixed Metal) was an alloy of rare earths of the following approx compo: Ce 49.0, La 25.6, Nd 16.0, Pr 4.6, Sm 2.0, Ti 1.0, Y 1.0, and Fe 0.8%. It was used as a component of delay elements for electric blasting caps. Other ingredients of delay elements included: Mg, Al, Ni and Zn homogeneously mixed with a fuel such as Si and an oxidizing agent such as PbO_2 .

Reference:

- H.M. Kerr, C.R. Hall, USP 2,560,432 (1951) C.A. 44, 1259 (1952).

Mischmetall (Mixed Charge). Designation for a mixture of lead oxide and lead sulphate for use in detonators. (See also Sprengkapsel A and Sprengkapsel B).

Reference:

- W. Schneider, Sprengtechnik, 1952, Bd 10/11, p. 196.

Mittel AEP (Agent AEP). Trade name for Ethyl Ester of p-Toluenesulfonic Acid, $H_3C \cdot C_6H_4 \cdot SO_3C_2H_5$; white crystals m.p. 31-32°, its solution in organic media was claimed to be a good gelatinizer for NC.

Reference:

- Karl-Metz, Chemische Untersuchung, Braunschweig (1944), p. 161.

Mittel KP (Agent KP). Trade name for Cetyl Ester of p-Toluenesulfonic Acid, $H_3C \cdot C_6H_4 \cdot SO_3C_{18}H_{37}$, brown oil d. 1.207 at 15°. Its sol in organic media was claimed to be a good gelatinizer for NC.

Reference: Karl-Metz, Chemische Untersuchung, (1944) p. 161.

Mollit II. German trade name for Central II.

Mollit II. German trade name for Central II.

Monochit (Monochite). According to Marshall (Ref 1) monochit were Favier type explosives. According to Colver (Ref 2) these explosives were invented by Kasl in Germany. Table 26 gives the composition of some monochits.

Table 26

Designation	An aluminum	K K nitrate	TXN TXN	Colloid Colloid	Char- coal	Alkali chloride
Monochit I	81	5	13	-	-	-
Monochit II	64	5	14	1	1	17
Monochit III	64	5	12	1	1	19

Abbreviations: TXN Trinitroxyloxy

According to Sterribach (Ref. 3), Monochit was an explosive suitable for loading projectiles and it was prepared by mixing ammonium nitrate with the solid and liquid products of distillation of solvent alcohols. (See also Filler No 37, under Fillers).

References:

- 1) Marshall v 1 (1917), p. 592
- 2) Colver (1918), pp. 238 & 234
- 3) Sterribach, Schieß- und Sprengstoffe (1933), p. 278.

Monobol. See general section.

Morser (Mörser). See under Weapons.

Morser Bomb. See under Bombs.

Morser Shell. See under Grenades and under Spigot Mortar Projectiles.

MP-14 (Solid Catalyst) used for decomposing the T-Spelt (hydrogen peroxide) in liquid rocket propellants.

Broken porcelain pieces, previously coated in a 50% solution of T-Spelt (q.v.) and dried at 110° for 24 hours were cooked for 10 minutes in a 50% solution of 2 parts Ca per manganate and 1 part K chromate and then redried at 110° for 24 hours.

When generating steam from T-Spelt, copper coils were used with MP-14 in order to accelerate initial decomposition. The rate of catalyst to copper was about 2 to 1.

Reference: CIOB Rept 30-115 (1945), p. 11.

M-Spelt Commercial methyl alcohol, sp. gr. 0.796, used as a component of some liquid rocket fuels, such as C-Spelt (CIOB 30-115, p. 10).

"Mulsipede". Same as Hochdruck Pumpe (High-Pressure Pump).

Munition. See Ammunition.

Munition-Pist Mine. See under Landmines.

Murphy's Changing Device. Used for linear adjustment of the range of some electrical time fuses, consisted of a cylinder, which fitted around the barrel of a cap just behind the muzzle and was connected by means of an electrical cable to a battery and a voltage-control mechanism located at the breech and of the gun. A charging ring, located in front of the muzzle, was held by means of three arms placed 120° apart. These arms also served for conducting the electric current from the cylinder to the ring. When a projectile equipped with an electrical time fuse, such as the Type 8/30 (EIZAS 5/30), reached the muzzle, the "fused wire" (located on the outside of the fuse and connected in

its storage time. This an electric range control mechanism was a step 550 volt. If the wire 16.00 seconds. No adjustment voltage applied. Reference: CIOB (1953), pp. 27 & 28.

Muzzle Protection in

Mysel (Mysel) of methyl

Mysel was

The material

was to be im-

and shock,

continuous

methanol and

new stable

described in

pp 9-10. For

more power

other high

comparable

mobile liquid

insoluble in

lean solu-

ical action

in methanol

directly in

that is be-

of the pure

mixture con-

methanol. T-

Mysel

new house

Mysel: Rosen-

155 methyl-

to H₂O

00 weight

methanol.

Follow-

mixture

to 7500-420

1/4 g. power and brisance-comparability to sheet-comparability to comparable to that of aliphatic. (PHTN). Like NC Mytol causes explosion, but they disappear more or coffee proved most successful. Explosives.

nitrate and its mixtures with benzene, etc found extensive ingredients of numerous liquid nates and explosives. Some of an Eisenzapferstoffe (substance

explosives or propellants, Mytol (poly) was used either by itself liquids, such as benzene, MNB or was replaced completely by case of plastic explosives or and right small amounts of NC the case of solid explosives used with a large amount (25-30%) of benzene mixed with the usual ones, such as kieselguhr, sawdust,

Mytol is a volatile liquid, all be kept in airtight containers. During plants were built in Ger. of WW II and the total capacity of ton per month. The largest plant was at Ebnick of Dynamit

used for the following purposes: were used as rocket propellants, torpedoes, land mines, bombs, hand grenades, fuses, etc. were used as military demolition charges. They were used as bursting charges in mines, hand grenades, torpedoes, etc. were used as military demolition charges. They were used as bursting charges in mines, hand grenades, torpedoes, etc.

in on Mytol Explosives and their use. When used for military or mines. When used for destroying or, etc. Mytol could be placed in, thus avoiding being of holes. present, they could be easily all demolition charges (such as fully on the surface of a rock, the underground wall, liquid Mytol be covered by pipes (also the end of the train derailed were found to be suitable for

such as methyl nitrate 75-80 and added to be satisfactory as a the rate of propagation in this the combustion chamber to the that this mixture could not be where in the range of 200 to

D) Liquid Mytol was found to be suitable for clearing out enemy trenches, foxholes, woods, etc. This clearing out operation was necessary "in order to destroy mines, or other explosives of toxic devices left by the enemy." The following ingenious method, using Mytol in the form of vapor, was developed by the Germans:

A bomb provided with two fuses, filled with Mytol and containing a small box with liquid carbon dioxide was dropped from a plane on the target. The impact of the bomb caused the first fuse to burst the box with CO₂ and to break the bomb. This caused the vaporization and distribution of the Mytol throughout the trench (or foxhole) without igniting or exploding it. The second fuse (time fuse) caused the detonation of the explosive mixture consisting of Mytol and an explosive oxygen. With sufficiently strong initiation the following reaction has been postulated:



When using this bomb in cold weather, the vapor pressure of the mixture can be increased by incorporating a small amount of methyl nitrite, CH₃ONO

E) Liquid Mytol, or straight methyl (or ethyl) nitrate, was used in the following device developed by Smeidgen:

Two small glass ampoules (bombs) one filled with methyl nitrate (or with less volatile ethyl nitrate) and the other with metallic sodium were placed inside a fuse closed on an HE filler of a land mine, but separated from it by a thin sheet of plastic material. On top of the bulb was placed a glass support. Pressure on the trigger caused crushing of the bulb. This was followed by an explosive reaction between methyl (or ethyl) nitrate and sodium. As a result of this, the sheet of plastic was pierced and the explosive charge inside the mine or bomb detonated. Based on this principle, several land mines were developed. The smallest and simplest land mine consisted of a flask containing 80-90 g of Mytol. Through the neck of the flask was inserted a short tube reaching nearly to the bottom of the flask. An ampoule containing metallic sodium was placed in the neck tube and on top of it a long plugger was carefully inserted. The pressure of this plugger caused bridging of the ampoule in the neck tube so that it was in contact with the Mytol. This action caused the detonation of the Mytol in the flask. The efficiency of these small mines was sufficient to disable a motor vehicle etc. Larger mines consisted of rectangular sheet-iron boxes filled with 25 g of 88/12 Methyl nitrate/MNB mixture and used the Mytol-sodium fuse.

F) Liquid Mytol explosives were also used to increase the penetrating effect of shaped charges, such as 40/60-TNT/RDX explosive. For this, a small glass ampoule (bulb) filled with 90/10-methyl nitrate/MNB mixture was placed in the air space (stand-off space) between the concrete surface of the shaped charge and the object to be pierced, such as armor, concrete, etc. For maximum effect the initiator (fuse) should be placed at the end of charge farthest from the target and pointing towards it. For instance, in shaped charge torpedoes, initiation of the explosive should be started from the tail end and not from the nose, as it is done in ordinary torpedoes.

G) Soft jellied explosives could be obtained by incorporating 3 to 5% of NC in any of the Mytol explosives, as for instance, the ones containing MNB. These jellies could be also mixed with pulverized solids, such as sodium nitrate and/or rock powder, thus obtaining solid explosives. The 40/60 mixtures were found suitable for filling the 50 kg projectile mines. These mines exerted a strong blast effect.

H) Hard jellied propellants could be prepared by incorporating in liquid Mytol (such as the base costs 75-80% of methyl nitrate and 20-25% of methanol, or MNB) comparatively large amounts (25-30%) of Autocollabols. Such mixtures formed very uniform hard colloids without pores or cracks and for this reason were found to be suitable as solid rocket propellants. It is believed that some of these mixtures were used toward the end of WW II as a fuel for V-1 and V-2 rockets.

Because of high volatility of Mytol, the propellant sticks used in rockets had to be coated with a special material impervious to Mytol.

1) A hard jellied explosive prepared by gelatinizing NC with a mixture of 91-95% methyl nitrate and 5-9% of MNB, was used in torpedoes.

2) A solid, highly bluish, explosive consisting of 30 to 40% of 75/25 Mytol mixed with such amounts of hydrated Ca nitrate and lignin that the oxygen balance was equal approximately to zero. The mixture was found suitable for filling bombs and land mines.

Notes: a) The high blueness and fairly high sensitivity to shock of the final mixture was presumed to be due to the fact that Ca nitrate entered and bound some methanol of the mixture, thus leaving part of methyl nitrate as free sensitive droplets. Another explanation of free methyl nitrate was partial evaporation of methanol, which in some explosive then methyl nitrate. According to Dr. H. Walter, Mytol's vapor in the form of an explosive mixture coming about 25% methanol b) In order to prevent an excessive liberation of free methyl nitrate, it was proposed to use a solvent less volatile than methanol such as benzene or nitrobenzene. In order to prepare such a mixture, the regular Mytol, which is a mixture of 75% methyl nitrate and 25% methanol, was shaken with benzene or MNB in presence of some water. This caused the methanol to go into the aqueous layer, while methyl nitrate remained mixed with benzene or MNB. c) A solid explosive consisting 30% of a mixture consisting of 50 parts of methyl nitrate and 10 pin of benzene, plus 55% of hydrated Ca nitrate, 10% of finely pulverized aluminum and 25% of pulverized, wet, and the oxygen balance equal approximately to zero. It was highly bluish and greenish although its nitrogen content was much lower than that of TNT (14.25 vs 36.5% for TNT). This mixture was proposed as a filler for warheads in rockets V-1 and V-2.

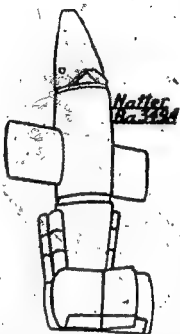
Notes: Mixtures of methyl nitrate 90% with benzene 10%, do not undergo very significant change in composition in storage. The composition of Mytol mixtures may be easily and rapidly determined by checking it's refractive index.

L) A solid Mytol explosive consisting of 65/15-Methyl nitrate/MNB gelatinized with NC and mixed with sodium and hydrated Ca nitrate was suitable for use in hand grenades or in mining.

M) A solid bluish explosive consisting of Mytol and a pulverized mixture of K nitrate, aluminum, and pent was suitable for hand grenades, land mines, and rock blasting. References:

- 1) G.R. Rees, Report on Explosives, PBL Rep. 65, 160 (1945)
- 2) H. Walter et al, German Development in High Explosives, PB Rep. 78,271 (1947)
- 3) J.C. Teichmüller, Chem. Ing. News 32, 2586 (1934) (Propellants for Rockets and Space Ships).
- 4) "Neubaus" (Nikolovsk). A self-propelled mine formerly known as the "Hohlraum" consisting of an 80 mm A/T gun on a Pafler IV or on a modified Pafler IV (See also under Pafler).

Motor B-349A. A surface-to-air, piloted missile developed in 1944 at the Bachem Werke GmbH. It was propelled by hydrogen peroxide/methanol + hydrazine hydrate and carried 35 R4M rocket projectiles in its stage. Launching weight 4800 lb, overall length 21.21 ft, width 58.0", max range 24.8 miles and max altitude 49,400 ft. It originated from a vertical ramp and climbed at a velocity of 35,300 ft/min. Reference: K.W. Gadand, Development of the Guided Missiles, "Flight" Publications, London (1952) pp 10 & 114-15.



Motor B-349B. A surface-to-air, piloted missile developed in 1945 at the Bachem Werke GmbH. It was propelled by hydrogen peroxide/methanol + hydrazine hydrate and carried in its nose 24 RZ 75 PBM (q) rocket projectiles. Launching weight 4,525 lbs, overall length 20.0 ft, width 16.0", and max altitude 50,000 ft. It was launched vertically and controlled by a radio link to the pilot in conjunction with ground control. Reference: K.W. Gadand, Development of the Guided Missiles, "Flight" Publications, London (1952) pp 114-5.

Nebeisäure (Fog-Acid) is a smoke-screen agent consisting of 30/50 - Chlorosulfonic acid/Sulfuric acid, (by weight) Reference: R.A. Richardson et al, C.I.O.S. Rep. 25-18 (1945), p.6.

Nebeiswerfer 41. See under Rocket Launchers.

Nebeiswerfermündung (Shunt-Circuit Igniter) is described in the book by Heyling-Greif (1956), p. 216.

Neofluor-Projektivile. See Arrowhead-Projectile.

Neofluor-Projektivile. See Arrowhead-Projectile and also German Projectile.

Neofluor. The name given after WW I to explosives used for rock blasting, spot-lighting stumps, etc. These explosives were prepared from a surplus military explosive called Hexanit, which consisted of certain hexanitrochlorophenylamine 60-70 and TNT 40-30%.

References:

P. Neesen, Schlegel and Sprengstoffe, Drogen (1927), p. 71.

100

Composition (%) and some properties	Nicro-baronite A	Nicro-baronite B
Aluminum	5.0	2.0
Am. astrate	82.0	69.0
Nicroglycerin	5.6	22.0
Collodion cotton	-	0.75
Liquid BNC	5.0	3.0
Petroleum tar	1.5	2.0
Wood meal	1.5	1.25
Ph Block Expansion (Picric acid = 100) (See "C" & "p" in the French Section.)	124.0	125.5

TABLE 27a

Composition (%) and dimensions	Nipolit tubes	Nipolit sticks
NC (12.6-12.7% N)	34.1	29.1
DEGN	30.0	20.0
PETN (unwaxed)	35.0	50.0
Stabilizer	0.75	0.75
MgO	0.05	0.05
Graphite	0.1	0.1
Length of grain	80 mm	30 mm
Diameter of grain	27 mm	9.1 mm
Hole Diameter	9.1 mm	—
Hole Depth	19 mm	—
Weight of grain	42 g	—
Caloric value, cal/g	1300	—

Note: MgO was added to neutralize acid developed on decomposition, and graphite was added to prevent the accumulation of hazardous static electrical charges.

For the preparation of Nipolit, a water slurry of NC was air-grated in a lead-lined vessel, with the desired amount of DEGN. After 15-20 minutes stirring the mass was centrifuged to remove all but about 25% of water and the resulting cake was treated, at about 50°C, in a Werner-Seidler machine with the calculated amount of pulverized PETN, some water, stabilizer, MgO and graphite. After about 15 minutes of tumbling the mass (plate) was transferred to rubber lined bags where it was allowed to age for 48-72 hours.

a) According to Ref 4, all raw materials with the exception of PETN were added in the paste mixing stage, while PETN was added during incorporation.

b) It was claimed that the aging process insured better gelatinization and reduced the tendency to fire during the rolling operation which followed.

c) The caloric value of the materials was carefully adjusted to between +30 and -10 calories as permissible variation from specification value for the propellant being processed. If outside these limits, the material was returned to the mixer and the caloric value either reduced by adding cellulose or hydrocellulose or increased by adding wet paste consisting of NC and DEGN. Each mixer was sampled at least every 8 hours. For a total charge of 10 kg a margin of 1 kg of report material was permitted.

Rolling and granulation were carried out as follows: About 18 kg of the aged paste was passed, about 15-20 times, through a pair of vertical rolls maintained at 90-100° (Ref 3).

Note: According to Ref 4 rolling was conducted at a temperature not higher than 75°C.

The resulting sheet (moisture content about 3%) was made by hand into a carpet roll and transferred to the press-house where it was kept in a steam heated oven, prior to transfer to the extrusion press. Then the mass was extruded at a pressure of 200 kg/cm² and at a temperature of about 80° and the resulting tubes (or sticks) cut into desired length.

After drying the cut material for about 24 hours at 40-50°, the moisture content was reduced to about 1%. The next operation consisted of writing each stick of Nipolit with acetone and pushing the stick into the tube of Nipolit flush with one end. This left a cavity 70 mm long in each tube to accommodate a detonator. The stick Nipolit (core) acted as a booster.

References:
1) O.W. Stickland et al, PB Report 1820 (1945), p 1b
2) A.A. Swanson & D.D. Sager, CIO Report 29-24 (about 1940), p 29

3) I. Litke, Prezentat Chemistry 27(4), 487-94 (1948)
C.A. 43, 4465 (1949) "Recent Developments in the Field of Explosives" (Translated by Dr Irvin Simon of Arthur D. Little Inc.)

4) A.A. Swanson, D.D. Sager & L.M. Sheldon, Ordnance Target Report No. 88 (Spec Rept No. 2073), Manufacture of Solventless Type Powder and Nipolit by the Deutsche Sprengbombe, Krieger Vks.

Nitric Acid (Salpeterminerale). Its preparation, properties and uses are described in the general section. Nitric acid was produced in Germany during WW II, mostly by the ammonia oxidation process, in quantities exceeding 140,000 tons per year. In addition, there was also available the 17,000 tons produced in occupied Austria, Czechoslovakia and Poland.

For the manufacture of highly concentrated (80% ammoniacal) nitric acid, the so-called "Hoho" (qv) process was developed.

Production of nitric acid in Germany was controlled by the Stickstoff-Syndikat.

Following is a partial list of the principal producers of nitric acid in the Western Zone of Germany:

- Badische Anilin- und Sodafabrik A-G, Oppau (formerly IG Farben AG)
- Bergwerksgesellschaft Hibernia, A-G, Hesse, Stickstoffwerke, Wanne-Eickel
- Chemische Fabrik Kalk GmbH, Köln-Kalk (Founded in 1877)
- Elektro-Nitrum A-G, Rhine, bei Launenburg (Baden)
- Fabrikwerk Höchst, bei Frankfurt a/Main (formerly IG Farben AG)
- Gewerkschaft Victor Chemische Werke, Cosmopol-Rhein 2, Werdau
- IG Farbenindustrie A-G with plants at Leverkusen (formerly Fried Bayer & Co., Bochum-Gerthe, Ruhr (called Chemische Werke Lebrungen GmbH) was founded in 1916) and Herford-Sodingen, Ruhr (formerly GAFAG)
- Ruhrchemie A-G, Oberrhausen-Holten, Ruhr (founded in 1927 under the name of Kohlenchemie A-G)
- Virtachische Forschungsbau GmbH (VFO) with plants at Embach, Kr. Lüneburg (founded in 1939-1940) and at Langelsheim, Hara (founded in 1939)

According to Ref 3 the following plants in the Eastern Zone were dismantled and shipped to Poland or Russia:

- Chimzavodsk a/d. Bolsh, Brandenburg (Dynamit A-G)
- Sinterfeld South (described in Ref 1)
- Elbsiedler
- Heydeck
- Leuna
- Passau (Bayerische Stickstoff A-G)
- Siedelwerke
- Witten (described in Ref 1)

References:
1) R. Mosley, BIOS Final Rept 889, June 22 (1946)
2) A. Krawczyk & P. B. B. BIOS Final Rept 1232, Item 22 & 31 (1946)
3) F. M. Davis et al, BIOS Final Rept 1442, Item 22 (1946).

Nitrobenzol (Nitrobenzol). An early type of aluminized explosive. The following mixtures, described by L. Néard, New Ann Fr 22, 596 (1948) are given in Table 28.

Table 28

Composition (%) and some properties	Nitrobenzol A	Nitrobenzol B
Aluminum	3.0	2.0
Am nitrate	82.0	59.0
Nitroglycerin	3.0	22.0
Cellulose acetate	—	0.75
Liquid DNG	5.0	3.0
Peroleum tar	1.5	2.0
Wood meal	1.5	1.25
PB Block Explosive (Picric acid = 100)	124.0	125.5

(See "C" in the French Section)

Nitrocellulose, Nitrocellulose oder Schießbaumwolle, abbreviated in German to Nz (Nitrocellulose, abbreviated in this work to NC). See general section under Cellulose. Due to the absence of saccharose in Germany, thus nitrocellulose was prepared from wood pulp.

Following is a brief description of the method used during WW II at the Krimmel Fabrik of D-A-G, as given in Ref 1 & 2:

- Bleached cellulose in the form of crêpe paper (made from wood pulp), was broken down in special machines into flocks and then blown into large drying chambers where the moisture content was reduced from 6-7% to 1-2%.
- 25 kg of cellulose flocks were fed with stirring into a nitration of 0.7 m³ capacity containing 1125 kg of mixed acid (MA), prep'd by fortifying the spent acid (SA) from previous batches.

Note: For NC of 11.25-11.50% N, called pe-Wolle, the MA consisted of 20% nitric, 62-64% sulfuric and 16-18% water; for NC of 13.2-13.5% N, called Schießwolle, the composition of MA was 22% nitric, 67.5-68.5% sulfuric and 9-10% water. The time of nitration was 30 minutes at the temperature 30°.

- The contents of the nitration were emptied into a centrifuge (one for every 4 nitrations) and spun for 6 minutes at 500 rpm.
- The separated spent acid (SA) went to rotating filter drums where the small lump particles of NC were separated and then to the dryer.

c) The NC which was removed from the centrifuges and the filters was carried by a stream of water into pressure washers where the bulk of the acid was removed by stirring with water.

- The slurry was then pumped to a preliminary boiling vessel provided with a double bottom of which the upper one was false, consisting of a screen through which the wash water was allowed to flow off at the end of the boiling period. Boiling was carried out at atmospheric pressure: 3 hours for PE-Wolle and 6-8 hours for Schießwolle.
- After removing the acid water, the NC was carried by a stream of water into the pressure boiling plant, where the material was cooked for 6 minutes in stainless steel autoclaves, starting at 100° and finishing at 142-145°.

Note: Pressure cooking had a double purpose: it reduced the viscosity of NC, to the desired level and it speeded up the stabilization. The details of the pressure cooking varied from plant to plant.

- A sample of cooked NC was sent to the laboratory and if the viscosity of the NC (as dictated by the stopper method in a 3% acetic acid) was within the desired range, the charge was dropped into a pulping machine such as the Hollander or Banning-Seybold. Here the NC was beaten for several hours, while the pH of the slurry was maintained between 7 and 9 by adding soda periodically. It usually required 3 to 4 kg of soda.
- The pulped NC plus water was pumped into vertical rotating sieves where more water was added. Here the smaller particles of NC passed through while the larger particles were retained by 90° mesh. The larger particles were removed by aspirators to be repulped, while the slurry of smaller particles went to a dewatering device (rotating drum sieve).

j) The dewatered small-particle material was transferred

to a final emulsifier consisting of a cylindrical vessel in which the NC was mixed with live steam until the slurry was brought to a boil. Then the water was decanted, the NC washed with water and a sample sent to the laboratory, in case of colloidal content (PE-Wolle), the above treatment was usually sufficient and the material would pass the Bergman-Jank Test (Hasting for 2 hours at 132° C should not produce more than 2 ml of NO per 1 g of PE-Wolle).

b) If the material was guanoform (Schneiswolle) the treatment was not sufficient and no testing had to be continued until a satisfactory B-J Test value was obtained (Not more than 2.5 cc NO per 1 g of Schneiswolle).

d) In order to obtain NC of the desired H content and viscosity, several batches were blended in large vats provided with stirrers. The blended material was then stirred in a large quantity of water and run through grit traps.

Notes: Grit traps were round pans, conical at the top. The slurry entered from below and its velocity decreased as it flowed upward (due to the increase in diameter of the vat) so that an extent that all the heavier particles (such as grit or dirt) dropped to the lower part of the vessel while the particles of NC continued to travel upward.

a) After "degrittling", the slurry was thickened up by passing it through a dewatering rotating drum for final dewatering. The partly dewatered material was sent to a centrifuge where it was spun at 1000 rpm. b) The remaining NC was shovelled into a zinc-lined line container (provided with a cover, where it was weighed, labelled and dispatched either to propellant plants or to a plant manufacturing "Kobolpervass" (Ray Photo).

c) As the waste water from the manufacture of NC contained an appreciable amount of suspended small particles of NC, it was required that these particles be removed before the water was allowed to leave the plant site. One method was to allow the water to run through so-called Danish traps. These were conical vessels with the narrow part at the bottom. The water flowed from the bottom upward; as the area of the vessel decreased, the velocity of flow was reduced to such an extent that the suspended particles settled. The accumulated fines were periodically removed from the vessel.

Notes: In near-German propellants were then examined at Picnic (Amsel) during WW II; the nitrogen content of the NC was around 127.02%, which means that the NC was intermediate between the PE-Wolle and Schneiswolle. One of the DGBN propellants contained NC with N+10.5% (See under Propellants).

Stutzhuber (Ref. 3) describes briefly various methods of manufacture of NC and gives compositions of mixed acids used for the preparation of NC with nitrogen contents of 11.6, 12.5, 12.75, 13.2, 13.4 and 13.7%. Yields and solubilities of various nitrocelluloses in 30% ethylalcohol mixture are also given.

References:

- 1) O.V. Shickland et al., General Summary of Explosive Plants, PR Rep 925 (1941), pp 50-53.
- 2) Lee Nutting et al., Manufacture of NC at the Krümmel Plant of the Dynamit A-G, PR Rep 16,666 (1945).
- 3) A. Stutzhuber, Spreng- und Schießstoffe, Rauscher, Zürich (1948) pp 62-66.

Nitrocellulosopropellant (Nitrocellulose Propellant or Single Base Propellant). See under Propellants.

Nitrocellulose. A low-tensile explosive oil used in the manufacture of some dynamites. It consisted of 80% di-nitrophenylol and 20% NG and was prepared by nitration of commercial amorpholohydria containing glycine (P. Naude, Schiess- und Sprengstoffe (1925), p 113).

Nitroform or Trinitromethane, described in the general section, was prepared and investigated during WW II in Germany, by Dr. Schimmelochmidt. He recommended the preparation of nitroform from nitro-acetone, potassium hydroxide and hydrogen peroxide, according to the following reaction:



His preference for the above method was based on the claim that the method previously employed by Onnes and McKay, depending on the reaction between trinitroacetone, K hydroxide and hydrazine, is hazardous since, in addition to K salt of nitroform, azido-acetic acid and azo nitroform (as was previously believed), is formed:

Nitroform was liberated from its K salt by distillation at reduced pressure in the presence of sulfuric acid.

The resulting product had a mp of 26.4° as against 22° obtained by some previous investigators.

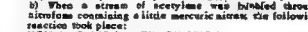
Dr. Schimmelochmidt also found that nitroform may be extracted from the reaction product of nitroform and sulfuric acid using liquid nitrogen dioxide, as in a solvent. This method of nitroform recovery was considered to be the greatest improvement, where the product so obtained could be converted to trinitroacetone with only a small amount of sulfuric acid. (See also under Trinitroacetone).

Notes: Due to the strength of sulfuric acid, which developed in Germany during WW II, any substance which could be used in place of sulfuric acid was considered highly desirable. For this reason, the use of liquid nitrogen dioxide was proposed also for the extraction of other nitrocompounds in addition to nitroform.

Nitroform was found to be an excellent mask inhibitor for the incorporation in polyvinyl acetate emulsions and also was found to be superior to NG nitroform in that it did not destroy the emulsion.

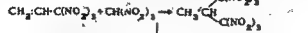
In the course of the investigation of the reactions between nitroform and organic compounds Dr. Schimmelochmidt observed several substances which were highly explosive at the reaction.

a) On treating nitroform with vinyl-methyl ketone, the following reaction took place:

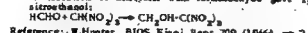


The resulting Trinitrovinylmethyl ketone was an explosive comparable in power to RDX.

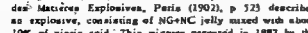
b) When a stream of acetylene was bubbled through nitroform containing a little mercuric nitrate, the following reaction took place:



Interaction of this compound with nitroform gave an extremely powerful explosive, believed to be a mixture of 1,4-Di(trinitro)butene and Trinitrobutene:



c) Reaction of nitroform with formaldehyde gave Trinitroethanol:



References: W. Hantz, BIOS Final Rep 709 (1946), pp 2 & 6.

Nitroformol peritum. Under this title J. Daniel, Dictionnaire des Matières Explosives, Paris (1902), p 323 described an explosive, consisting of NG+NC jelly mixed with about 10% of picric acid. This mixture, patented in 1887 by the Deutsche Sprengstoff Gesellschaft of Hamburg, did not prove to be very stable.

Nitroglycerin and Nitroglycol (Nitroglycerin and Nitroglycol, abbreviated in this work to NG and NGG). The manufacture and properties of these substances are described in the general section under Glycerin and Glycol, respectively.

In Germany the nitration of glycerin or of glycol (ethylenglycol) was conducted either by a batch process or by a continuous method, such as that of Schmid, Meissner or Bianzi. The nitration was made either separately for glycerin and glycol, or more often as mixtures, such as glycerin 60 and glycol 40%.

The batch method of nitration of glycerin, or of glycol or of their mixtures at the Krümmel Fabrik Dynamit A-G may be given as an example:

a) 300 kg of glycerin was run into 1470 kg of mixed acid, consisting of HNO₃ 50, H₂O 52 and H₂O 25, acid, contained in a stainless steel nitration vat which was provided with an air stirrer and cooling coils.

b) In order to maintain the mixture in the nitration vat at about room temperature, the brine, cooled to as low as -12° was circulated through coil.

c) After about 25 minutes of nitration, the air agitation was stopped and the mixture allowed to stand, in order to accelerate the separation of the organic production.

d) 2 g of an 80/20 mixture of Na fluoride and of ignited kieselgur was added.

e) The separated oil was air-dried at 12° with 400 liters of water and after removal of the water, the oil was air-dried for 12 minutes at 40° with 500 l of 2% soda ash solution.

f) After cooling the mixture to 28°, while still containing to stir, 50 g of pulverized calc was added and then the mass was allowed to stand.

g) The separated oil was run through a pipe which coded some distance short of the storage tank. From that end of the pipe, the oil was transported to the tank by means of hand trucks.

h) The spent acid which in the case of NG weighed about 1200 kg and had the approx. comp: HNO₃ 7.5, H₂O 75 and H₂O 19.5% and in the case of NGG 15% (nitrocellulose) weighed about 1030 kg and had the approx. comp: HNO₃ 8.5, H₂O 74.5 and H₂O 17%, from which the bulk of oil had been removed, was allowed to stand for 60 days in lead-lined vessels, called "Alfenzapfentanks".

The separated oil was washed in a small auxiliary vessel first with water and then with soda ash solution.

Notes: The total yield of oil was reported to be about 233 parts per 100 of glycerin. Other plants reported yields ranging from 25 to 274, and for NGG 230.

h) The spent acid of (g), was blown by compressed air to a tank and from there to a separator in order to recover some more of the explosive oil. Then the acid was transferred to the Recovery Plant where the nitric acid was distilled off, leaving weak sulfuric acid as a residue.

i) As the waste wash waters of operations (d) and (g) contained small amounts of oils (NG, or NGG) it was necessary to remove the oils before allowing the waters to run into a stream, lake, etc. This was accomplished by allowing the waters to run through large settling tanks, sometimes installed in cascade form.

j) In order to economize on the consumption of nitric acid to prevent poisoning of personnel all nitric acid fumes (as well as nitrogen oxide gases) were drawn from both the nitration and separator by means

of a suction device and led to an absorption tower in which they were met by a spray of water to dissolve them and form nitric and nitrous acids.

k) A sample of washed oil [see operations (d) and (g)] was sent to the laboratory for testing. The Abel test was usually about 40 minutes.

Note: The results of the Abel Test were usually higher than in the US practice. The high German results are presumably due to the fact that calcium was used in the separation of the oil [see operation (e)]. The Americans do not use calcium to improve the separation of NG or of NGG from spent acids.

The Stryken Tank of B. S. A. G. also used the batch process, while the Schiebhaus Fabrik of Dynamit A-G had three different NG installations:

a) Batch plant.

b) Continuous plant with Meissner nitration and Diazal separators and washers.

c) Continuous plant with Bixtal nitration, separator and washers, installed by Meissner, Bixtal, Switzerland.

In the Bixtal installation, which had an output of 800 to 1000 kg per hour, the nitration was a cylindrical stainless steel vessel approximately 2 ft in diameter by 8 ft 6 in deep (See Fig. 1, A/29 of Ref 5). Cooling was carried by means of cooling coils through a series of concentric coils suspended inside the nitration vessel.

Stirring was carried out with a mechanical stirrer situated in the center of the inner cooling coil and running at a speed of 400 rpm. A tangential separator was placed about 2 ft below the level of the outlet of the nitration and a 2nd separator followed the 1st. The mixed acid used in the nitration was approximately 30/50-nitric acid/sulfuric acid, stored in a tank for at least 10 days and then passed through a stainless steel pipe before use.

Procedure:

a) The mixed acid, 3 parts, and glycerin (or glycol, or glycerin plus glycol) 1 part, each metered by means of a counter, entered continuously and simultaneously.

b) The emulsion consisting of nitrated product (oil) and spent acid left the nitration and was run straight to a tangential separator placed about 2 ft below the level of the outlet from the nitration.

c) The separated acid/oil went to a stainless steel vessel 1 1/2 ft in diameter and 2 ft deep, provided with a mechanical stirrer, where the oil was washed with an equal volume of water, while the spent acid (which in case of NG had the approximate composition: HNO₃ 11, H₂O 87.6 and NG 0.75%) went to another separator.

d) The acid/oil mixture was then washed with a special lead separator, called Scheider. This operation permitted the removal of some additional oil from the acid was forced to be reused for nitration of the next batch, or before the acid was sent to the recovery plant.

e) After pre-washing the oil with water, the emulsion flowed continuously into a tangential separator from which the separated oil went to the next part of the process.

f) The acidic water (which in the case of NG had the approximate composition: HNO₃ 10.6, H₂O 89.4, H₂O 8.6 and NG 0.75%) went to another separator.

g) The acid/oil mixture was then washed with a special lead separator, called Scheider. This operation permitted the removal of some additional oil from the acid was forced to be reused for nitration of the next batch, or before the acid was sent to the recovery plant.

h) After pre-washing the oil with water, the emulsion flowed continuously into a tangential separator from which the separated oil went to the next part of the process.

i) The acidic water (which in the case of NG had the approximate composition: HNO₃ 10.6, H₂O 89.4, H₂O 8.6 and NG 0.75%) went to another separator.

j) The acid/oil mixture was then washed with a special lead separator, called Scheider. This operation permitted the removal of some additional oil from the acid was forced to be reused for nitration of the next batch, or before the acid was sent to the recovery plant.

k) The pre-washed oil of the operation (d) went through two vessels in series, each of them equipped with a stirrer. Simultaneously with the oil at 75% soda ash

solution, measured by a micrometer, entered the vessels. There was no separation of the emulsion between the vessels, and the oil/soda emulsion went from there to an annex (wash-house), located outside the mound surrounding the siting-house.

Note: All the above listed operations were conducted in the siting house. It should be mentioned that the siting was provided on the bottom with a glass plate which could be broken when it was required to drown a charge. A pneumatic hammer operated as a handle at the side of the building was used for breaking the glass. The drowning tank, located below the siting, contained about 5 times the volume of the siting of 95% sulfuric acid.

b) The emulsion from the previous operation went through two separators located in the wash house. The separator oil was collected in a rubber lined aluminum truck holding 600 kg, while the wash waters went via a cascade system to a tunnel leading to the Rhine River b) The truck conveyed solidified oil was emptied into a storage tank where it was allowed to stay for at least one day to permit the water to separate.

Note: In a sewer type of final settling house, there were 6 flamm-magnesian lead-separators placed in cascade and working continuously.

1) The dried oil was removed from the storage tank as needed, by means of heavy rubber, buckets of 40 kg capacity.

The average yield of dry NG from the Blaas plant was 232 parts by weight per 100 parts of dry glycol. The stability was 14 minutes by the Abel Test at 15°. When the nitroglucose was made from acids recovered from TNT manufacture, it was sometimes necessary (in order to obtain satisfactory quality for NG) to incinerate from 0.1 to 0.2% of the acid in the acid washing liquor. During the war, however, diphenylamine stabilizers were sometimes used when the quality of the NG was unsatisfactory. In the manufacture of double-base propellants, NG was used alone, while in the manufacture of commercial dynamite-type explosives it was used in mixtures with NG (nitroglycerol).

- References:
- 1) R. Schaefer, Nitroglycerin und Dynamit, Veit, Leipzig, (1906)
 - 2) P. Nozin, Nitroglycerin und Nitroglycerin Explosives, Williams & Wilkins, Baltimore, 1928) pp 25-178 & 210-235
 - 3) A. Stettbacher, Schaefer und Sprengstoffe, Barth, Leipzig, (1933), pp 146-177
 - 4) O. W. Stickland et al., General Summary of Explosive Plants, PB Ref 925 (1945), pp 67-6
 - 5) R. A. Schaefer et al., Investigation of German Commercial Explosives, B I O S, Ref 833, Item 5, H M S O, London (1946), pp A-1/4 and A-2/4
 - 6) A. Stettbacher, Spreng- und Schießstoffe, Zürich, (1948), pp 59-62

Nitroglycerin-Nitrocellulose Explosives: Commercial explosives suitable for blasting rocks were prepared by mixing double-base propellants (left as surplus after the formation of WW II) with other ingredients, such as inorganic nitrates and organic nitrocompounds. Following are the compositions of some of these explosives:

- a) Mining List No 33 Explosive: NG 50 to 40, NC 60 to 70, with added 0 to 5% of hydrocarbons of oil (and/or naphthalene) and 0 to 10% of paraffin (and/or urethane, and/or centralite, and/or dicyanamide)

- b) Mining List No 35 Explosive: NG/NC jelly 94 to 96 and 4 to 5% of a 50% aqueous solution of Ca nitrate
- c) Mining List No 16 Explosive: NG/NC jelly 97 to 99, and 1 to 3% of substituted urethane.

References:

- 1) P. Nozin, Nitroglycerin etc., Baltimore (1928), pp 449-50
- 2) J. P. Lefebvre, Poudres, Paris (1933), p 458.

Nitroglycerin-nitrocellulose (Nitroglycerin Explosive). See Dynamit.

Nitroglycerin-polymer (Nitroglycerin Propellant). A propellant based on NG and NG, also called double-base propellant. Prep and properties of typical NG propellants are given in the book of A. Stettbacher, Spreng- und Schießstoffe, Zürich (1948), pp 41-43

See also under Propellants.

Nitroethyl (Nitroethyl), abbreviated in this book to NGE is described in the general section under Glycol. The manufacture of NGE in Germany was conducted in the same manner as for NG. Because of high volatility, it is not advisable to use NGE alone in explosive compositions (although this sometimes did), but it is satisfactory to add NGE to NG in order to improve the burning point of the latter. Such mixtures were used extensively in the preparation of commercial dynamite-type explosives. References: Same as under Nitroglycerin.

Nitroguanidine (Nig) [Nitroguanidine (NGu)], described in the general section under Guanidine, was prep'd in Germany by treating guanidine nitrate (GuN) with concentrated sulfuric acid as described by Schaefer (Ref 4).

Briefly, the method was as follows: In order to obtain 100 kg of NGu, 135 kg of GuN was added gradually to 300 kg of 98% sulfuric acid while stirring and cooling so that the temperature was not allowed to go above 45°. The resulting mixture was run into a dilution vessel (maintained at 0°) in which the precipitation of the crude NGu took place. By using a centrifuge, the crude product was separated from the liquid phase, which contained about 20% H₂SO₄. The crude material was dissolved in boiling water, mixed with the mother liquor from the previous batch (see below), made strictly neutral by means of ammonia, filtered, and the filtrate cooled to at least 45° at low pressure. The resulting crystalline suspension was transferred by air pressure to a centrifuge. This gave a purified NGu with a water content of about 6% and a mother liquor which was later used for dissolving the crude NGu of the next batch (see above) (Ref 4). The preparation of NGu was also described by Stettbacher (Ref 1).

Uses of NGu:

- A) According to Davis (Ref 7), NGu in admixture with Am nitrate and/or paraffin was used during WW I for lighting various bombs. These compositions were highly insensitive to shock.
- B) During WW II NGu was used either in propellant such as the cool, aromaticless and flashless triple-base propellant, called Gudopulver, or in explosive compositions.

Note: When intended to be used in propellants, the NGu were required to be of such size and shape that the ingredients of a propellant were rolled into sheets, the

incorporation was smooth and rapid. When intended for use in explosives, two kinds of NGu crystals were used:

- a) finest grain crystals (dust) obtained by rapidly evaporating a hot aqueous solution of NGu under high vacuum. These crystals were found to be suitable for pre-ignition.
- b) crystals with high bulk density (above 1.0), obtained by crystallizing NGu in the presence of collodion. Such crystals were found to be suitable for the casting of TNT-NGu mixtures.

C) As an example of the use of NGu as a high explosive may be cited the 1800 kg AP bomb in which some NGu was placed in the nose as a sort of protection (bumper) for the more sensitive main charge consisting of "Filler 109".

Note: According to CIOB Ref 32-18 (1945), German production of NGu towards the end of WW II was about 1500 tons month.

- References:
- 1) A. Stettbacher, Nitrocellulose 7, 141-145 (1936) (Nitroguanidine)
 - 2) T. L. Davis, Army Ordnance 26, 93 (1939)
 - 3) PB Ref 925 (1945), pp 24 & 216
 - 4) W. Schaefer, PB Ref 16-603 (1945)
 - 5) Allied & Enemy Explosives, Aberdeen Proving Ground, (1946), p 149
 - 6) A. Stettbacher, Spreng- und Schießstoffe (1948), p 44.

Nitroisobutylglycerolnitrate (Nitroisobutylglycerol Trinitrate). See general section and also A. Stettbacher, Spreng- und Schießstoffe, Zürich (1948), p 69.

Nitrool. See general section.

Nitrool. An amol type explosive in which TNANs (trinitroammonol) was used to replace TNT. The mixture of TNANs 60 and Am nitrate 40% was of light yellow color with a m.p. about 75° which permitted cast-loading. Its strength, brisance, and sensitivity to mechanical action were similar to those of 40/60 Ammol. It was hygroscopic and in the presence of moisture the TNANs hydrolyzed to picric acid, which would attack metals with the formation of dangerous picrates, while the Am nitrate could hydrolyze to form ammonia. Nitrool was used in some sea mines and torpedoes.

References:

- 1) Allied & Enemy Explosives, Aberdeen Proving Ground, Md., (1946), pp 110-11.

Nitropropellants. German research on the preparation and properties of nitropropellants is described in CIOB Refs 35-41 (1945). See also general section under Propellants.

Nitropropene (Np). See Pentrit (PETN).

Nitropropenolnitrate. See Pentrit.

Nitroquin (Nitroquin). See general section under Starch.

Nitroretol. See general section under Toluenes.

Nitrous Oxide, N₂O. Same as Gm-I. See also general section.

Nitroxyl. See general section under Xylene.

Nitrocellulose. Same as Nitrocellulose.

Nitroguanidine (Nitroguanidine). See general section under Sgns.

Nitrool. See under Starch Section.

Nobelit (Nobelit). A type of permissible gelatin-dynamites was made before and after WW I. Two examples are given in Table 29.

Table 29

Composition (%) and some properties	Nobelit	Nobelit 19
NG (gelatinized with NC)	28.7	26.0
Dextrin	2.5	2.0
Wood meal	1.0	1.0
Potato flour	10.0	-
Vegetable oil	0.3	-
Am nitrate	39.7	34.8
Am chloride	17.6	12.0
Saturated soda of Ca nitrate	-	5.0
Oxygen Balance, %	-	5.0
Density	-	1.75
Velocity of Detonation, m/sec.	-	3750
Transit Time, sec.	270	220

(See also Water-Nobelit)

- References:
- 1) P. Nozin, Schieß- und Sprengstoffe (1927), p 150
 - 2) P. Nozin, Nitroglycerin (1928), p 407.

Nobel's Sprengöl oder Sprengöl. Same as Nitroglycerin.

Nobel's Vetterlymann. One of the older permissible dynamites: NG 50, Am nitrate 51, flour 39, wood meal 6, naphthalene 2, and alum 1%. Veloc of detonation 3850 to 3930 m/sec at 1.16 (Marshall 2 (1917), p 492).

Non-Destructive Testing of Materials. Some of the German methods of testing are described in B I O S Final Rep 609 (1946). See also general section.

Normal Gas Volume (Normal Gas Volume). Volume of gas at normal temperature (0° or 20° C) and normal pressure (760 mm Hg) or of volume with NTP. Calculation of the volume of gas developed on explosion is described in the general section.

(See also A. Stettbacher, Spreng- und Schießstoffe, Zürich (1948), pp 13-14.)

NSP. See under Ignition.

NalmenNP. See under Ignition.

Oberflächenbehandlung (Surface Treatment). See general section under Surface Treatment of Explosives, Propellants, Pyrotechnic Compositions, etc.

Oberon. Given: A device designed in 1944 for controlling the burning point of the aircraft incendiary rocket, R 104 BS. It was claimed that the Oberon device improved the chance of a rocket from a negligible value to a probability of about 0.6.

Reference: T N 5985-2 (1953), p 255.

Oberring Bullet, caliber 7.22 mm, developed by the Deutsche Waffen- und Munitionsfabrik A-G, Laback, exploded with a flash on hitting the target. The bullet consisted of a steel casing containing a charge of white phosphorus, a detonator and a striker with a steel spring. The base of the casing was closed with a lead plug. Reference: H. P. Pelot et al., CIOB Ref 33-20 (1946), pp 26-7 (See drawing on next page).

A unique light and effective AP projectile was designed for use in the Russian 76.2 mm A/T gun. At first the Germans attempted to adopt the arrow-head type projectile Page 40 but found it unsuitable. In its place they developed a projectile of normal shell design, but employed a plastic laminar armor to give body to the shell and still keep it relatively light. This shell, described by E. Engemann in the *Ordensburg* (Sept. 1944), p. 112, consisted of the following components: a) a ballistic cap of an aluminum alloy, screwed onto the shell, b) an armor-piercing core, consisting of tungsten carbide with a central steel core, pressed into a steel core holder, c) a sleeve of molded plastic surrounding the core and its holder and filling the space between the body and these components, forming an armor head like the ballistic cap. The plastic had a fairly high shock resistance.

Still more effective was Arrow of handle Type projectiles designed by G. Gessner.

The projectile constructed at the Röchling Plant at Saarbrücken were very effective for penetrating concrete. (See also under Arrow Projectiles, Arrowhead Projectiles, German, German Projectiles, Röchling Projectiles and Sabot-Projectiles).

Panzerschnecke. See under *Hautschilde*.

Panzerschneckenmine. See under Landmines and also p 262 of TM 9-1985-2 (1953).

Panzerschreck. *Panzerschreck*, *Panzerschreck* and *Pippen* were the shaped charge weapons developed before and during WW II in Germany.

The *Panzerschreck* was the shaped charge rocket, similar to the American Bazooka, but was heavier and had a shorter range than the latter. It was developed by Dr. Panzerkretz, which was a better weapon with a range of 150 meters. Another weapon, called the *Pippen*, was essentially the L.S. on *Panzerschreck* mounted on a light carriage. The *Panzerschnecke* was a long-range weapon for shooting a shaped charge, developed by the Rüstwerke Co. It was a rocket-bore 8.0 cm mortar. (See also under 80 mm and 80 mm Weapons).

References: 1) L.E. Simon, *German Research in WW II*, Wiley, N.Y. (1947), pp 167-8
2) A. Scherhauer, Spreng- und Schmelzstoffe, Zürich (1948), p. 154.

Panzerschreck 42. See under Rocket Launchers.

Panzerschreckmine. See under *Panzerschreck*.

Panzerschnecke (L). A shaped charge head grenade, introduced by the Luftwaffe for use in close combat against armor vehicles of all types. Diameter of body 11", overall length 21", weight 2.1 lb. It was provided with four collapsing fins. The fins were folded against the handle. When the grenade was thrown, the fins spring open and stabilized the projectile in flight.

References: 1) A.J. Derr, *Ordance Sergeant*, Oct 1945, p. 8
2) A. Scherhauer, *Ordance Sergeant*, No 7 (1945).

PANZERWURFMINEN (L)

Papmine. See under Landmines and also on p 261 of TM 9-1985-2 (1953).

Perschute Flare. See under *Flare*.

Pewmen. Mixture of Am perchlorate 90 and peroxide 10% used for military purposes. (A. Scherhauer, Spreng- und Schmelzstoffe, Zürich (1948), p. 91).

Pflichter Bombing. A night bombing tactic developed during WW II in Great Britain used against the German. The tactic consisted of dropping bombs on a target previously illuminated by flares dropped from the leading planes.

This method permitted more accurate bombing of the target. Reference: A. Scherhauer, *Pflichter Bombing*, (1953).

(See also *Pflichter Bombing* and *Pflichter Bombing*).

Pewmen. See *Cartridge*.

PC 1400 PX was a mule controlled glider bomb, released from aircraft and designed for attack against rail and smaller objects. [TM 9-1985-2 (1953), p. 195-6].

Pewmen. A rocket-researcher, including an air tunnel, constructed in 1956-1957 in the isolated spot on the German Baltic coast. The first rocket developed at Pewmen was the A-4, the predecessor of the A-4 rocket, commonly known V-2.

A fairly detailed description of *Pewmen*, Röchling and its activities is given in Ref. 6.

Pewmen is now in the Eastern Zone of Germany.

References: 1) A. Scherhauer, *Les Amers Service*, Allendamm, Paris (1947), pp 103-110
2) L.E. Simon, *German Research in WW II*, Wiley, N.Y. (1947), pp 33 & 110
3) J. G. Tschalitz, *Chem Eng News* 22, 2382 (1954)
4) W. Dörmer, V-2, Vitik, N.Y. (1954).

Pewmen. See also *Pewmen* (PETN).

Pewmen. See *Swiss* section.

Pewmen or *Pewmen* corresponds to the American *Pewmen*, described in the general section. (See also *Pewmen* No 16, 17, 28, 42 and under *Pewmen*).

Pewmen. See under *Swiss* Explosives.

Pewmen or *Pewmen* (NP). See general section under *Pewmen* or *Pewmen* (NP). It was manufactured in Germany by hand, commonly in small quantities.

A) The batch method was essentially the same as that used in the U.S.A.
B) The continuous method, as conducted at *Pewmen*, Fabrik, D.A.G. consisted essentially of the following operations:

- Nitric acid of the highest concentration and PE in the ratio of 5 to 1 were introduced simultaneously into a mixture of 50 liter capacity. The PE was added by means of a "driving" machine feeding at the rate of 600 g every 47 seconds. The temperature was maintained at 15-20° by means of cooling coils.
- The solution-suspension of PETN in nitric acid was led to an stir mixer, where the mixture was maintained at 12°.
- After this it went to a third vessel, where a strong jet of water diluted the acid and precipitated that part of PETN, which was dissolved in the stronger acid.
- The slurry was run through a vacuum filter and the per was rinsed several times with water.

e) The precipitate was transferred to a vessel where it was heated in dilute soda ash solution to 80-85° and from which PETN was run on a 2nd filter.

f) After separating the liquor by vacuum filtration, the PETN was washed with water and separated to a moisture content of 1-10%.

g) The moist material was dissolved in 98% acetone preheated to 50°, and allowed to run gradually and with stirring into a vessel containing cold water.
h) The acetone was distilled off and the crystallized PETN separated from the bulk of the water by vacuum. It was then packed in rubber bags and carried to the packaging house.

i) For phlegmatization (desensitizing) PETN, the *Pewmen*, Fabrik, D.A.G. used either *Monna Wax* or a synthetic IG Wax - 41a. The amount of wax added to PETN was usually 10%, although mixtures with as high as 60% were known. The crystals of PETN were suspended in cold water containing some common salt solution. The temperature was raised to about 40° and molten wax was added, in a thin stream. The temperature was raised and the mass maintained at the boiling point until about 70% of the water had evaporated.

j) The slurry was then cooled (by adding cold water and filtered). After washing the phlegmatized product with water and removing as much water as possible by suction, the product was dried to reduce the moisture content to below 0.1%. The material was then screened and packed.

k) The semi-continuous method as practiced at the *Kübel Fabrik*, D.A.G. was essentially as follows:

l) The nitric acid apparatus consisted of 3 stainless steel vessels connected in series. A charge of 200 kg of PE and 1000 kg of 99% nitric acid was fed into the first reactor (which was cooled with coils circulated in coils and in a jacket) where the main nitric acid place at 15-20° during about 10-15 minutes. A second charge of PE and HNO₃ was meanwhile weighed and transferred to the first reactor immediately after the 1st batch was transferred to the 2nd reactor (which was also provided with cooling). Following this, the 1st batch was transferred to the 3rd reactor, the 2nd batch to the 2nd reactor, and a 3rd charge was introduced into the 1st reactor, etc. The total time of nitration was about 40 minutes.

m) In the 3rd reactor, the mixture was diluted with water to give a waste acid of about 30% strength.

n) After filtering off PETN from waste acid, PETN was washed with water and then digested with soda ash solution in a neutralizing vessel at 60° until the slurry was weakly alkaline (pH about 10). This was followed by water washing directly on the filters.

o) The next operation, crystallization from acetone, was done in a continuous manner in a battery of 6 distillation vessels connected in series. In these vessels, water was added to the solution and the acetone gradually evaporated leaving a water slurry of PETN. After removing the bulk of the water by vacuum filtration, the moist PETN (10% H₂O) was transferred to the next phlegmatization.

p) Phlegmatization was carried out in a water slurry of 315 kg of PETN (containing 10% H₂O) plus 1200 kg of water at 85°, to which was usually *Monna Wax* or IG Wax - 41a, in the proportion of 1 part wax to 9 parts PETN by dry weight, was added with stirring.

Notes: According to German Railroad regulations, phlegmatized PETN was permitted to be shipped if contained at least 6% wax. Unphlegmatized PETN required at least 30% of water for shipping.

q) PETN was also phlegmatized by the addition of TNT (10 to 30%) and the operation was conducted by suspending PETN in about 6 parts of water at 70° heating to about 80° and adding molten TNT with agitation. This was followed by cooling, filtering and drying. The mixture was allowed to be shipped dry (Ref. 1).

r) The manufacture of phlegmatized PETN at the *Wollfhausen Plant* was described by Swanson Ref. 3 and CLOS Ref. 25-16 (1945).

s) Abbreviation: PE Pentacetylene trioxide.

t) References: Same as under *Pewmen* and *Pewmen*.

Pewmen or *Pewmen* (Pewmen) Explosives. *Pewmen* (Pewmen) was used under the name of *Filler No 3-NP* as a bursting charge in some grenades and small shells (such as the 20 to 30 mm), as well as in a lower detonator. *Pewmen* was also used as a phlegmatized shell. (Ref. 1).

The use of PETN desensitized with 10% wax was much more common.

Note: The wax used in German explosives was usually *Monna Wax*, obtained from the lignites found in many parts of Germany and consisting of about 75% of the wax. The properties of *Monna Wax* are comparable to those of *Canabax* was imported from Brazil. German PETN-wax mixtures were usually deep pink. The explosive properties of such mixtures were the same as those of the corresponding American mixtures described under *Pewmen* or *Pewmen* in the general section.

The principal uses of PETN-wax mixtures were as follows: fillers for various shells, bombs, grenades, and some air mines; fillers in some shaped charges; standard boosters in chemical and incendiary ammunition; standard sub-booster in all types of ammunition and as the core in a detonating fuse.

Explosives desensitized with TNT, are briefly described under *Pewmen* or *Pewmen* as well as *Filler No 16, 18, 28, 32, 34, 36, 37, 42* etc. in some mixtures of PETN were used in underwater ammunition.

Besides these mixtures there was also a plastic explosive (see *Filler No 43*) and explosives consisting of PETN, RDX and wax (see *Filler No 45*).

References:

- Amor, Alfred and Enay Explosives, Aberdeen Proving Ground (1945), pp 38-42
- O.V. Sackman et al., General Summary of Explosive Plants, P.B. Rept No 925 (1945), pp 42-45
- A.A. Swanson et al., Manufacture of Phlegmatized PETN, P.B. Rept No 320 (1945)
- A. Scherhauer, Spreng- und Schmelzstoffe, Zürich (1948), pp 66-67.

Pewmen or *Pewmen* (Pewmen). See general section and also under *Pewmen*.

Pewmen Explosives. See *Pewmen* or *Pewmen*.

Pewmen or *Pewmen* (Pewmen). A type of industrial explosive based on perchlorates. Table 31 gives some perchlorates listed in the book of *Nachrichten* (Ref. 1).

Table 31

Ingredients:	Composition, %		
	1	2	3
K perchlorate, of which up to 10% of the total explosive may be replaced with Am nitrate and/or K nitrate	50-75	62-73	-
K and/or Am perchlorate Am nitrate	-	-	30-40 15-45
Note: When Am perchlorate is incorporated most of the Am nitrate is replaced by K nitrate in an amount chemically equivalent to the amount of Am perchlorate.			
Vegetable meal	1-5	-	5-8
Vegetable meal and/or solid hydrocarbon	-	1-4	5-8
Nitroglycerin (angelcatalized)	3-6	-	-
Nitrocellulose of cellulose and/or acetoneless and/or diphenylamine in which up to 4% of the total explosivity may be substituted with nitrocellulose	20-50	20-30	15-20

Seutcher (Ref 2) lists the following perchlorates:

Table 32

Composition, %	Perchlorates		
	1	2	3
K perchlorate	68	35	34
Am nitrate	10	42	48
TNT and DNT	-	14	-
DNT	16	-	12
Wood (or vegetable) meal	1	5	6
NG (nitroglycerin)	4	41	-
MNN (mononitrophenylene)	1	-	-

References:

- 1) P. Nozom, Nitroglycerin, Berlin (1928), p 431
- 2) A. Seutcher, Schiene- und Sprengstoffe, Leipzig (1935), p 316.

Perchlorat-Sprengstoff (Perchlorate Explosive or French Matter). According to P. Nozom, Schiene- und Sprengstoffe (1937), p 133, the following explosive mixture, developed during WW I at Zeuzenstille für wissenschaftlich-technische Untersuchungen in Neußhagen, was found to be suitable for use in Verfahrtschächten (trench meters): K perchlorate 56, DNT 32 and DNN 12%.

Note: This explosive was called Perdit by Davis (1943), p 364, but on p 118 Nozom gives a different formulation for Perdit.

Perchlorat-Sprengstoffe (Perchlorate Explosives). Explosives based on the perchlorates of ammonium, potassium or sodium were used to a limited extent in Germany, as for instance: Permanno, Perchlorite, Perchlorit, Perdit, Perchloronit, etc.

(See also Perchlorate Explosives in the general section).

Note: According to Davis p 364 the perchlorates recovered from surplus bombs etc. of WW I (see Perchlorat-Sprengstoffe and also Perdit) were used in the German post

WW I commercial explosives, such as Perchlorit, Perchlorin, Perchloronit and Perchlorit. When the supply of surplus perchlorates became exhausted the manufacture of perchlorate explosives was entirely discontinued because the price of new perchlorate was too high.

Perchlorit (Perchlorin). A type of perchlorate explosive used in mining before and during WW I. Table 33 gives the composition of two perchlorites.

Table 33

Ingredients and properties	Composition, %		
	1	2	3
K perchlorate	35	34	-
Am nitrate	42	48	-
DNT	10	0	0
DNN	4	0	0
Wood meal	3	0	0
Coal powder	-	2	-
NG	4	-	-
Oxygen Balance, %	+1.7	+1.7	-
Tensile Test, cc	340	340	-

*DNT was prep. by the reaction of m-DNT.

Reference: Nozom, Nitroglycerin, Berlin (1928), p 133.

Permanno (Permannoite). A blasting explosive which replaced Carosil in such quarries and one mine: K perchlorate 65, NG 5, aromatic nitrocompounds 25 and vegetable meal 5%.

Reference: J. Behr, Manual of Explosives, Macmillan, N.Y. (1943), p 116.

Perdit (Perditin). An explosive developed during WW I as a replacement for the Corps of Engineers Explosive, (Pinnemont) Donait. The composition and properties of Perdit were: Am nitrate 72, K perchlorate 19, wood meal 5 and a mixture of DNT and TNT 1% (density 1.20-1.25, Tensile test value 370-580cc, sensitivity to initiation: required at least a No 5 cap for detonation).

It was used not only as a demolition charge but also for loading bombs and trench mortar shells.

References:

- 1) P. Nozom, Schiene- und Sprengstoffe, Dresden (1927), p 118
 - 2) A. Seutcher, Schiene- und Sprengstoffe, Leipzig (1935), p 309.
- (See Note under Perchlorat-Sprengstoffe).

Perchlorit (Perchloronit). A type of mining explosive manufactured after WW I from K perchlorate recovered from surplus military explosives. Table 34 gives two examples. (See next page).

Permanno (Permannoite). A type of mining explosive manufactured before WW I by the Sprengstoff A-G Carlsberg. One such explosive, called Gerstein-Permanno, was described in this section under Gerstein-Sprengstoff. Table 15 gives two examples of Permanno.

(See next page).

Table 34
(Perchlorites)

Composition and Properties	1	2
	1	2
K perchlorate	58	59
Am nitrate	8	10
DNT + TNT + vegetable meal	30	31
NG (nitroglycerin)	4	-
Oxygen Balance, %	+2.2	+1.8
Density	1.53	1.52
Velocity of Detonation, m/sec	5000	4400
Tensile Test, cc	340	350
Ph Black Crawling, mm	20.0	18.0
Regulate for initiation minimum	No 3 cap	No 3 cap
Cap Test, cc	6.0	4.0
Heat of Explosion, kcal/kg	1170	1160
Temperature of Explosion, °C	5145	5115

References:

- 1) P. Nozom, Nitroglycerin, cc, Berlin (1928), p 430
- 2) T. L. Davis, Chemistry of Powder and Explosives, Wiley, N.Y. (1941), pp 364-5.

Table 35
(Permanno)

Composition and Properties	1	2
	1	2
K perchlorate	52.5	51-54
Am nitrate	42.3	59-63
NG	-	3-4
Cellulose cotton	-	1-1.5
TNT	16.0	11-13
Sand	12.0	5-9
Wood meal	3.0	1.5-3.5
Mixture	-	0-2.3
Veloc of Detonation, m/sec	3780	-
Density	1.13	-
Tensile Test, cc	-	305
Cap Test, cc	-	8.0
Impact Sensitivity (2lb weight)	-	20 cm

Permanno were used in ponds and in two mines. Some permanno were on the British Permitted List and on the Belgian SGP List.

Reference: A. Marshall, Explosives, London, v 1 (1917), p 384 and v 2 (1917), p 493.

Permanno (Permannoite). One of the perchlorate mining explosives manufactured from leftover stocks of WW I military explosives. The name Permanno is mentioned in P. Nozom, Schiene- und Sprengstoffe (1927), p 126, but the composition is not given.

Permanno (Permannoite). One of the pre-WW I explosives used in ponds mines and stone quarries: No nitrate 69, K nitrate 5, white 10, capri salt 15 and K bichromate 15, Tensile Test value 157 cc (vs black powder 108) and sensitivity to impact with a 2 kg wt 100 cm (black powder 65).

References:

- 1) A. Marshall, Explosives, London, 1 (1917), p 39
- 2) A. Seutcher, Schiene- und Sprengstoffe, Leipzig (1935), p 111.

PETN. See Perdit.

P.E.-Wells-A nitrocellulose of 11.25-11.50% nitrogen content, used for the manufacture of some smokeless propellants. See Nitrocelluloses and also Propellants.

Pfeilgeschoss. See Arrow Projectile.

Phenanthrene (Phenanthrene) was proposed by Römer to be used as one of the ingredients in explosives based on cyclotrimethylene-trinitrosamine (R-Salt), such as: R-Salt 96.5, phenanthrene 2.5, and DPhA 1.0%.

Reference: G. Römer, Report on Explosives, PBL Rept No 85, 160 (1946), pp 10-13.

Phenix Sprengstoffe (Phenix Explosives) were mining explosives patented in 1899 by the Sprengstoffwerke Dr. Nohmann & Co in Hamburg.

Table 36 gives some examples

Table 36

Ingredients	Composition %				
	1	2	3	4	5
NG	23	25	30	50	30
K nitrate	34	30	30	50	30
Am nitrate	1	35	32	30	32
Sandstone	40	-	38	-	-
Flux Flour	-	40	-	40	38

References:

- 1) Daniel, Dictionnaire des Matières Explosives, Paris, (1902), p 449
- 2) L. Gody, Traité des Matières Explosives, Neufchâteau, 1907, p 715.

Phenol (Phenol). See general section and also BROS Final Rept 1246 (1946).

Phosphorus Bombs. Some incendiary bombs contained phosphorus. For instance, the 50 kg Bred C50B bomb contained white phosphorus whereas the 50 kg Bred C50A bomb was filled with 50 lb of a mixture containing phosphorus 4, between 50 and pure rubber.

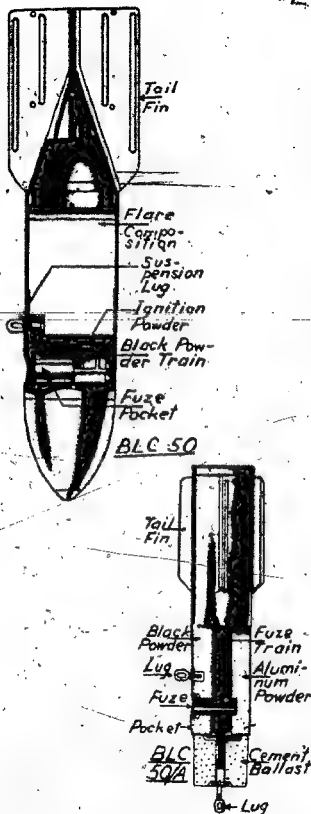
Reference: TM 9-1985-2 (1933), pp 14-5.

Phosphorus Grenade. One of the incendiary grenades made during WW II in Germany was described in BROS Final Rept 1253 (1946), p 2. It weighed 190 g and was prepared from a casing weighing 50 g, having a diameter of 105 mm. After filling the casing with a mixture of 50% wood meal (2 g) and aquaplastine (500 g), the 2 g was exhausted and the mouth of the grenade was inserted into molten yellow phosphorus. This operation allowed about 750 g of phosphorus to enter the grenade and impregnate the cotton and naphtaline.

Phosphor Bomb (Blitzschmelzende Bombe, abbreviated as BLC or BIC), called also Photographic Flash Bomb. German bombs were similar in external appearance to conventional 50 kg bombs and parachute flare cases. Their fillings, which could be either flare compositions or incendiary mixtures, were ignited by electrical or mechanical aerial burst fuses.

Following are examples of photoflash bombs:

- 1) BLC-50 bomb weighed 50 lb and resembled in appearance the SC kg Type I bomb except that the case was made of sheet steel with a heavy nose section. Body diameter 7.5", body length 26.4" and overall length 43.0" (see also under Bombs)
- 2) BLC-50/100 bomb consisted of a light steel casing 42.9" long and 8" in diameter. Its nose was filled with



concrete which acted as a ballast to stabilize the flight of the bomb. The outer section of the bomb contained 15 kg of Al Pyroshockil (qv), while the inner tube contained 3.5 kg of black powder, called Marine-Geschütz Pulver. This served for expelling, accelerating and igniting the Al powder, which continued to burn in the air. The black powder was exploded by means of an 80 mm long detonating fuse placed inside the tube passing through the black powder charge. The fuse was initiated by a small section of electric fuse inserted in the nose tail in the side of the bomb. Total weight of the bomb was 42 kg. The bomb was insensitive to bullet impact.

Note: The Pyrochloff aluminum could be replaced with an atomized Al powder called Griess, or by mixtures containing magnesium powder described under Photoflash Compositions.

Reference: TM 9-282-2 (1955), pp 63 & 81-3.

Photoflash Compositions. Among the compositions used by the Germans, may be mentioned the ones used in the ELC 50/A bombs.

a) 15 kg of flaked aluminum, called Pyroscorff (q v). It was insensitive to bullet impact and had the following characteristics: peak light intensity 490 million Hefner candles, time to reach peak intensity 70 milliseconds and total light output 63 million international candle seconds.

b) 30 kg of atomized aluminum, called Gotsen (q.v.). It was insensitive to bullet impact and had a peak light intensity of 800 million Rother candles. The time to reach peak intensity and the total duration of the flash were longer than for the 15 kg Pyrochiff.

c) 28 kg of pellets (13 mm diam and 7 mm high) composed of magnesium powder 39, Ba nitrate 33, synthetic phenolic resin, 6 and talcum 2%. It was sensitive to rifle bullet impact. Its peak intensity was 80% of that of Pyrochloff, and the time to reach peak intensity was 300 milliseconds.

d) 28 single-perforated pellets (60 mm diam and 220 mm high), each weighing 900 g (total weight of pellet 25.2 kg) and consisting of: Mg powder 50, Na nitrate 45 and wax 5%. A length of detaching fuse was passed through each pellet, and the ends of the fuse bound together.

It was sensitive to rifle bullet impact and had a peak intensity (measured through a yellow filter) 20% greater than for 15 kg of Pyroschliff. The time to reach peak intensity was the same as for Pyroschliff, but the duration of flash much longer.

Reference: TM 9-1095-2 (1953), on 82-4.

PH-Sals. (PH-salicyl). German name for Ethylenediamine-*N,N'*-bis(2-hydroxyphenyl)sulfone (EDDS), described in the general section. In Germany PH-Sals was prepd by treating ethylenedichloride with ammonia and NaOH, followed by amination with nitric acid not stronger than 50%. Although PH-Sals has a high $m.p.$ of 118°C, it is not the property of depressing the $m.p.$ of other explosives, completely soluble in water, and is not so sensitive that it no longer contains explosive mixtures. For instance, a mixture of 45% PH-Sals and 55% Am nitrate melts at 105° and can be cast-solid. Such a mixture has an explosive power equal to that of TNT or Ametal, but it has the "disadvantage" of shrinking "considerably" on cooling. Addition of aqueous Ca nitrate to this mixture practically prevents the shrinkage, and the mixture is a good cast. The following mixtures containing PH-Sals were used for filling some shells as a substitute for TNT.

n) Ammoniz: NH_4NO_3 46, PH-Salz 46 and $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ (tech) 8%; density of frugum 39-40 m. (See Fragments Density Test) ~

b) Ammonia: NH_4NO_3 55, PH-Salt 10, $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ 10, RDX 20, and NaN_3 5%; d 1.53, casting temp. 100°C , density of frozen mass 40 = (2.04 g).

c) H-5 (Ammonit): PH-Salz 10, NH₄NO₃ 50, KNO₃ 5, Ca(NO₃)₂·4H₂O 15, and RDX 20% (Ref 2)

d) S-16: PH-Gala 10, NH_4NO_3 , 32, NaNO_3 , 6 or 8, KNO_3 ,

2 or 0, RDX 10 and Al (powder) 40% (Ref 2)
e) 5-22 (Hexa): PH-Sala 14, NH₄NO₃ 45, NaNO₂ 9,
KNO₃ 3, RDX 14, and Al (powder) 15% (Ref 2)
f) 5-22 (Hexa): PH-Sala 14, NH₄NO₃ 45, NaNO₂ 9,
KNO₃ 3, HMDSPA 14, and Al (powder) 15% (Ref 2)
g) Amalot 41: NH₄NO₃ 52, PH-Sala 30, Ca(NO₃)₂·
4H₂O 6, RDX 10, and Menton wax 2% (Ref 3).

Compositions containing Al were particularly suitable for underarm weapons because they possessed high blast effect. Pz-Sel could also be used straight or slightly phlegmated. In the latter case, it was particularly suitable for use in anticoncrete shells, called Be-Granate (Be is the abbreviation for Betonconcrete).

References:

- 1) PB Rept No 925 (1945), p 24
- 2) PB Rept No 1820 (1945), p 29
- 3) PDL Rept No 89 160 (1946), p 25.

Plastic Acid - See **Pikric Acid**.

Planting or Penetration Test. For this test an explosive enclosed in an iron tube, 30 mm in diameter and 100 mm long with walls 3.5 mm thick, was detonated horizontally against a lead sheet 30 mm thick with sides 100 mm long. The penetration produced was compared with that of a standard explosive such as TNT.

References: G. Römer, *PBL-Rep* 35/60 (1966), p. 10.

Pfhe (Heckler) Missile. An experimental guided missile developed in 1941 by the Rheinmetall-Borsig Co.
Reference: K.V.Gatland, Development of the Guided Missile, London, (1952), pp 116-17.

Phthalene (Picric Acid) (P.A.). Methods of preparation and properties are given in the general section. It would be of interest to know that in 1892 the Chemiker Fabrik, Gröbenheim, Ger. Pat. 69 937, described a unique process for loading HMX shells of P.A. This was carried out as follows: A solution of P.A. and 5% of TNT was placed in a suitable container and the temperature was raised to a temperature of about 87°C which is slightly above the m.p. of TNT. On cooling there was formed a solid block consisting of crystals of P.A. cemented to the other intermediate layers of solid TNT, in place of TNT over solid TNT. This was done with not too high a m.p. may be used (such as TNT, DNP, etc.) and the same process may be used for any polymer. It was claimed that the resulting cemented block had high density, were easy to prepare, and were adapted for use in a mechanical action device.

During WW II P A was manufactured for use as a booster (compressed), as well as a filler for some shells, land mines, depth charges (see Filler No 2) and as a filler in stick hand grenades (see Filler No 3).

Case P A was used under the name of Filler No 24. Abbreviations: DNB Dinitrobenzene; DNCs Dinitroresol; DNPd Dinitrophenol; DNT Dinitroamine; YNCs Trinitroresol and TNT Trinitrotoluene.

References

- 1) E. de W. Colver, High Explosives, Van Nostrand, N Y, (1918), pp 319-20 & 697
- 2) Ames, Allied and Enemy Explosives, Aberdeen Proving Ground, Md. (1946)
- 3) A. Soerhachter, Spreng- und Schienstoffe, Rasthof, Zülich (1948), pp 75-77.

Place: San Silver.

Pistol (Pistole). See under Weapons.

Pistolpulver (Pistol Propellant). The following composition is given in Brunswick, Das rauchlose Pulver, (1926) p 136: gun cotton 90, Ba nitrate 1, DPhA 1.5, residual volatile gelatinizer 0.5 and moisture 1%.

Pistol Grenades (Pistolengranaten). Several types of German grenades were fired from special pistols, such as the 27 mm Walther signal pistol, etc.

(a) Pistol Grenade (Wurkgrat) [sic Leuchtpistole, 56].
consisted of a cylindrical egg band grenade attached to a
plastic stem (body) by a retaining tube. The plastic stem
contained the firing pin, delay igniter, detonator and a
base adapter for the propellant. The end of the stem was
closed before firing by a cardboard cap. After arming the
grenade, the cap was removed and the stem was inserted into the
target.

spread by withdrawing the safety pin, the plastic stem was placed in a barrel reinforcing tube which was previously placed in the barrel of the 27 mm Valther signal pistol. The cap and the propellant in the rear section of the stem were fired and the grenade went forward in the barrel.

range 80 yds). The impact of the grenade caused the firing pin to strike the primer and the resulting flash ignited (through the flash cubes) the delay igniter. After a delay of about 4.5 sec the grenade exploded (pp 340-1)

b) 27 mm Egg Type Pistol Grenade, described on pp 341-2, was fired from the latest type 27 mm Walther signal pistol, without the insertion of a rifled liner (as a reinforcing tube) in the barrel. The grenade was similar to the type 361 except in the construction of the stem.

c) 26 mm Pistol Grenade (Wurfgranatpatrone), fired from the smooth-bore pistol, 326 Leuchtpistole, consisted of a projectile having the appearance of a small mortar shell. A brass cartridge case, containing about 0.1 ounce of

the propellant, was crimped over the rear section of the grenade where the fins were located. The projectile itself consisted of an outer casing (body) and a loosely inserted inner casing containing the detonator and the main charge. The fixed fins are held by a cross-rod in the

charge. The firing pin, held by a coo spring, was located in the nose section of the body. The inner case was prevented from moving forward before firing by two metal balls fitting into a hole in the tail section of the projectile and resting in grooves. An arming (safety) rod

lized between the balls holding them apart. The withdrawal of the rod, caused by the setback on firing the projectile, allowed the retaining balls to move towards the muzzle thus releasing the rear section of the inner case. The case would now be free to move forward if it was not held by

would now be free to move forward if it was not held by the tension of the creep spring. This tension was overcome on impact thus allowing the detonator (contained in the inner case) to move forward and strike the fixed firing pin (pp 342-3)

d) 27 mm HE Grenade (Sprengpatrone) for the rifle pistol (Kampfpistole) consisted of a die cast aluminum body provided on the outside with five grooves making one quarter turn of the projectile. Inside the body was a steel cylinder containing two PETN/wax

cylinder containing 100 cc of water, pellets separated by a cardboard disc. The nose section contained the direct action fuse fitted with a protruding 'striker' head. The striker was held away from the fuse primer by 6 steel balls which rested in the groove of the striker and on a

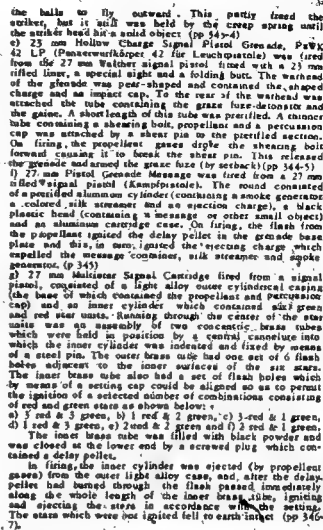
platform of the fuse. The balls were kept in position by a steel collar which was supported on three aluminum pins. A creep spring was located between the striker and the primer, and beneath the primer was an aluminum gaine containing in the upper part a plunger, bent and

containing in the upper part a mixture of lead azide and lead styphnate and in the lower part pressed PETN. Between the gainer and the main filling there was an air space. The propellant charge was contained in a cup which was placed in the cartridge attached to the tail section of the grenade.

There were 10 holes in the cup to lead the propellant gases to the base of the grenade. On firing, the gases propelled the grenade and retained it because of the rifling. The setback caused the collar in the fuse to move back crushing

the aluminum pins and the centrifugal force caused

GRENADÉS



Plastics in German Ordnance. During WW II there was a growing use of plastics in plants which manufactured acids, explosives and ammunition. For instance, linings for tanks and pumps, funnels, pipes, plastic trays for drying explosives, sealing-plugs in delay detonators etc. were made from plastic material. One of the plastics developed in Germany was Mipolam. Others were Novolac, Lignolite, Isopolmer, Trolicol etc.

Reference PB Rep't No 925 (1945), pp 7 and 25.

Plastic A plastic explosive of WW II: RDX 64, cotton 3.3 and liquid or semi-liquid* nitrohydrocarbons 32.5%. It was less efficient than the American Composition C2 because it contained less RDX. [Allied and Enemy Explosives, Aberdeen Proving Ground (1946), p 127.]

Plaster, Cellulose, Cellulose oder Camphresol Trade names: for p-Toluenesulfonamide, $\text{CH}_3\text{C}_6\text{H}_4\text{SO}_2\text{NH}_2$, white flakes, mp 137°, obtained as a by-product of saccharine manufacture. Its 20% alcoholic solution gelatinizes cold cotton completely at 55°.

[Kaoi-Hers, Chemische Untersuchungen, Braunschweig (1944), p 163].

Plastoments (Plastomenite). According to Daniel (Ref 1) plastomenites were propellants invented about 1889 by Gütler. They consisted of mixtures of the nitrated products of cellulose, sugar, starch, aromatic compounds, etc. with oxidizing substances such as inorganic nitrates, chlorates, chromates, etc. These compositions were modified beginning 1897 by incorporating 0.5 to 10% of colophony.

Colver (Ref 3) stated that Plastomene was a German propellant prepared by blending 5 parts of molten DNT with one part of nitrofin and sometimes small amounts of Ba nitrate. After incorporation the fused mass was granulated.

References:

- 1) Daniel, Dictionnaire des Matières Explosives, Paris (1902), p 634
- 2) A. Marshall, Explosives, 3, (1932), p 98
- 3) E. Colver, High Explosives (1918), p 169
- 4) H. Bruuswig, Das sauchlose Pulver (1926), p 134.

Plasticity. See Plasticity.

Plasticity or Plastit. According to Colver, High Explosives (1918), p 249, plasticity are plastic explosives patented by C.E.Bichel in 1906 (Ger P 181 574). They were prepd by mixing 85 to 87 parts of TNT with liquid or solid resins, such as copaliba balsam, benzin gum or styraz, with or without liquid DNT. The plasticity could be increased by incorporating some colled cotton. Table 37 gives some examples.
(See next page).

Plattenberchuss (Plate Shooting). A plate test for the estimation of the brisance of explosives similar to the one described in the general section.

Plastic Explosives. Several plastic explosives based on PETN and RDX were used in Germany during WW II. One of the earlier compositions consisted of RDX treated with "American vaseline" (see Note) until this vaseline became saturated. Therapeutic mixtures called Plastic and Hexaplast, which did not contain vaseline, were used. "American vaseline" was considered most suitable because it is "long life" and can be stretched like dough to form threads. European vaseline, such as the Russian, is not tacky and does not produce good plastic explosives. (See also Plastic Explosives in the general section).
Reference: PR Rep 925 (1941), p 74 & 77.

Plastics. (Kunststoffe, Pressstoffe): Manufacture and properties of plastics are described in the following:

- 1. W. Knauss, *Kunststoffe (in technischen Korrosionsschütz, Lehmann, München-Berlin 1945)*
- 2. H.S. Betgen, *Plast Report 1972 (1943)*
- 3. Anon, *Plast Report 19186 (1945)*
- 4. BIOS Field Reports: 282, 453, 445, 926, 1191, 1246 and 1729 (After IV)
- 5. BIOS Miscellaneous Reports: 1, 85, 67 and 98 (After VW II)
- 6. CROS Reports: 29-62, 32-26 and 33-23 (After VW II)

H. Sackring, *Handbuch der Kunststoffe (Fischer Buchverlag, München (1952))*

Table 37

Ingredients	Composition %			
	1	2	3	4
THT	87.0	85.0	87.0	85.0
Cetane-80	12.0	-	-	-
Laurel turpentine	-	14.0	-	-
Liquid styrene	-	-	4.3	-
Neosin gum	-	-	-	4.3
Liquid DMT	-	-	10.0	10.0
Colored cotton	1.0	1.0	0.3	0.3

Plasticizer-polymer (Black Cartridge Propellant). The following composition is given in Braunschweig, Das technische Pulver (1928), p. 136: cotton cord 23, gunpowder 74, diphenylamine 0.5, castor oil 0.5, nitrocellulose 1.0 and residual volatile plasticizers 1.0%.

PAP - Same as Pällipal 109 (Fr 109), described under Piller.

POL (Primer) also Löseemittel (Solventless Propellant) See under Propellant.

Pollapan One of the plastic materials developed prior to WW II by the Dynamit A.G., at Troisdorf. It is a non-toxic, non-flammable, non-explosive product.

References:
1) K. Reich, Kautschuk in Technische Kautschukmischungen, Lehmann, Jena (1943), p. 21
2) H. Schilling, in W. Ziemann, Kunststoffe-Taschenbuch, Chemie, München (1932), pp. 240 & 241
3) H. A. Tsch, Plastics, Arsenal; primer communication.

Polybort - See POL

Polybort. According to CIGS 21-3 (1945), a Nylon type polybort was developed at the Troisdorf Plant of Dynamit A.G. No description of its make and properties is given.

Polyglycol (PGK) (Polyglycol). A liquid product consisting of about 77% diethylene glycol (DEG), called in Germany Diylol, and 25% ethylene glycol (EG) called Glykol (1928). This product was made before and during WW II by IG Farbenindustrie starting with ethylene glycol in turn was obtained either from blast furnace gas (by liquefaction and subsequent fractionation) or by hydrogenation of acetylene. This means that no food materials were required for its manufacture. Known for the manufacture of plastic "critical food synthetics" such as fats were required. When this product was simulated, a liquid explosive was obtained which proved to be a better plasticizer for NC than NC. Another advantage of nitrocellulose (NPG) was that it produced much cooler propellants (possessing lower caloric value) than was ever possible with NC. See "G" Pulver.

Reference: G.W. Stickland et al, General Summary of Explosive Plastics, P.B. Rept No 925 (1945), p. 13.

Polyphenyl. A plastic composition which when applied to the surfaces of combustible solids prevented them from burning. It was used for coating the non-burning surface of solid rocket propellants.

Reference: TM 9-1965-2 (1953), p. 201.

Polyterrene Plastics. According to CIGS 21-3 (1945), p. 5, the IG Farben at Ludwigshafen produced two types of polyterrene which softened at 60° and 70° respectively. The copolymers of styrene were used.

Polyterrene Plastics. Proprietary and properties are described in CIGS Report 29-12 (1945).

Polyvinylchloride Plastic, called Lutivon, was manufacturer for injection molding because of its high melting point (over 200°). Considerable pressure was required to mold it and this caused rapid wear of the mold.

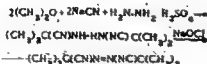
Polyvinyl Chloride (PVC) was used in Germany for the preparation of various plastics (Ref 1) and in most synthetic compositions (Ref 2).

The following polyvinyl chloride plastics are mentioned in Ref 1:

- a) Vinidur (q.v.)
- b) Moplen (q.v.)
- c) After-chlorinated PVC. It contained up to 60% of PVC and was very stable. It dissolved in methylene chloride in which the original PVC was insoluble.

Reference: 1) M.F. Taylor - F.J. Carro, CIGS Rept 21-3 (1945), p. 2
2) T. Ullmann, Encyclopedia, 37 (4), 487 (1944).

Powder N. Code number for the product prep by IG Farbenindustrie by condensation of acetone with sodium cyanide and hydrazine sulfate, followed by treatment with sodium hydroxide:



The product was used in the manufacture of powder materials such as from rubber space. It was used for Schottel tubes and submarine pipelines (see under Piller). It has the property of absorbing nitrogen when heated together with vinyl chloride in an autoclave at 150°.

Similar properties were displayed by Powder D (Diammoniumdinitrate) and Powder 254 (proprietary to Polyvinyl Chloride cyclodextrane instead of acetone). Reference: CIGS Report 21-18 (1945), p. 30.

Powder 254 or Stick Hand Grenade (Stielhandgranate) consisted of a wooden stick (handle) to which was attached a metallic can filled with an explosive. A similar type was the Japanese Type 98 Stick Hand Grenade and also the Russian Stick Hand Grenade.

Reference: TM 9-1965-2 (1953), pp. 319-320 (Stielhandgranate 24, 32 and 43).

Powder Identification. See Pulvermetallurgie.

Pre-synthesized Projectile. See general section.

Pre-synthesized Projectile. See Rifled Projectile.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-synthesized Explosives. German procedure is briefly described under Krimmel Pulver. Dynamit A.G., Pre-synthesized Explosives.

Pre-mixing (Primary Charge) is a top charge of a blasting cap or detonator. Explosives used for primary charges are described under Primary and Initiating Compositions.

Primary and Initiating Compositions. The following German terms are used in connection with this subject: Zündung (Primer Charge), Zündschicht (Primer Cap), Zündmittel (Initiating Composition), Zündständer (Initiator). A general

description of primary and initiating explosives may be found in Refs 1, 2, and 8 as well as in the general section. Refs 4, 5, 6, 7, and 8 are listed explosives used during WW I, but only in a few types of primers during WW II. Table 38 lists some German primary and initiating compositions used in fuzes, primers and detonators.

Table 38

		Composition %											Uses
	M.F	L.A	L.St	Sh.S	Temper	NC	Ca silicide	Oxidizer	Absorber				
a	85	-	-	23.5	-	-	-	KClO ₃	43.0	Glass 10	-	Primers in shells and some both fuzes	
b	-	-	94	-	-	6 (*)	-	-	-	-	-	Electric fuse primers	
c	-	-	88.7	-	-	11.3 (*)	-	-	-	-	-	Same as above	
d	-	-	37.5	7.4	4.2	-	12.4	Ba(NO ₃) ₂	38.5	-	-	Primers	
e	-	-	49.1	-	-	-	15.4	Ba(NO ₃) ₂	35.5	-	-	Primers	
f	-	-	32.1	-	-	-	-	Ba(NO ₃) ₂	47.9	-	-	Primers	
g	82	-	-	7	-	-	-	-	-	Glass 11	-	Primer-detonators	
h	60	-	40	-	-	-	-	-	-	-	-	Standard detonators	
i	55	-	45	-	-	-	-	-	-	-	-	Same as above	
j	14.4	-	85.6	-	-	-	-	-	-	-	-	Detonators	

* In compositions (b) and (c) the NC was made into a paste using any acetone. Then the paste was loaded to the ignition bridge of a primer.

Table 39 lists some cartridge case primer compositions used during WW II

Table 40

		Composition %					
		L.St.	Sh.S.	NC	APHV		
a	85	30 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	20 mm HE Shell	
b	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	57 mm APHV, 37 mm APHV, 47 mm APHV, 50 mm HE, 50 mm APC LC, 50 mm APHV, 50 mm APC SC, 80 mm CM and 88 mm AP Shell; some Lead Mire	
c	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	47 mm AP and 75 mm APC; 45 mm HE Shell	
d	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	47 mm HE Shell and 100 mm HE Shell	
e	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	47 mm HE Shell	
f	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	75 mm HE Shell	
g	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	75 mm HE and 105 mm HE	
h	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	88 mm AP Shell	
i	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm Mortar Shell	
j	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	Lead Mire (Tellemire 35)	
k	85	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	50 mm HE, 50 mm APHV, 75 mm HE and 75 mm APC	50 mm APHV, 75 mm HE and 75 mm APC	Lead Mire (Tellemire 42 and 43)	

Table 40 lists some primer compositions used in fuzes during WW II

Abbreviations: AP Armor-piercing; APC Armor-piercing, capped; CM Chemical mortar; HE High explosive; NC Nitrocellulose; PETN Pentanitroethyl tetraazide; R.N. Road mine; SC Self-decapating; L.A. Lead anode; L.C. Long

case; L.St. Lead anode; M.B. Monoblock; M.F. Mercaptan; MTF Mechanical time fuse; NC Nitrocellulose; PETN Pentanitroethyl tetraazide; R.N. Road mine; SC Self-decapating; L.A. Lead anode; L.C. Long

Table 42
Single Base (Nitrocellulose)
Propellants of WW II

Form	Composition, %						Other Ingredients	Uses
	NC	NN in NC	DPLA	Case	Acac	Graph		
Square	95.1	13.2	-	-	1.8	-	Unac	7.65 mm Mousier
Square	95.2	13.0	-	-	0.3	-	Unac	4.3 7.92 mm AP
SP	94.3	12.2	-	0.2	-	0.3	PETN	63.8 7.92 mm AP
							Unac	1.4
Square	95.1	13.1	-	-	1.0	1.0	Unac	7.92 mm AP
Square	95.0	13.2	-	-	-	-	Et carbamate	3.6 7.92 mm Ball, 7.92 mm Semi-AP, 7.92 mm AP and 7.92 mm HE
							K sulfates	
SP	92	12.5	-	0.4	-	0.6	PETN	66.0 7.92 mm HVAP
							Unac	7.0
Square	98.4	13.1	0.9	-	-	-	Unac	6.7 7.92 mm Rifle
							Graphitized	Grain A/T
SP	99.3	13.0	0.5	-	-	-	PETN	7.92/13 mm AP
SP	96.0	13.2	-	Same	-	-	Unac	34.0 7.92 mm HVAP
							Case & DNT	10.0
Square	96.1	12.7	-	2.6	-	0.3	Unac	7.92 mm AP
SP	95.0	12.2	-	2.0	-	0.1	Unac	-2.9 7.65 mm Mousier
							Unac	Pistol, 9.0 mm
	95.0	13.1	-	-	1.7	-	Unac	Pistol and 28/20 mm APHV
SP	96.4	13.0	-	-	0.3	-	Unac	3.3 9.0 mm Pistol
							Unac	2.1 9.0 mm Ball
							Unac	2.7 9.0 mm Ball, 9.0 mm Pistol and 70 mm Trench Mortar
SP	95.0	12.9	0.5	2.0	-	0.4	Unac	2.1 13.0 mm AP and 13.0 mm HE
SP	95.7	13.3	-	1.95	-	0.25	Camphor	0.95 20 mm AP
							Unac	3.15
SP	94.7	13.1	0.5	-	-	0.5	DBuPh	6.1 20 mm HE Mousier
							Unac	4.6
Square	99.7	15.2	0.5	3.4	-	0.5	K sulfates	0.5 20 mm HE Mousier
							Unac	1.8
SP	95.3	15.1	-	2.8	-	1.5	K sulfates	1.1 20 mm Inc
							Unac	0.5
SP	95.3	15.1	0.2	1.2	-	0.5	K sulfates	1.0 20 mm Solochon
							Unac	4.0
SP	94.1	13.0	0.4	2.4	-	0.4	Unac	2.7 13.0 mm AP, 13.0 mm HE, 15.0 mm HE and 28/20 mm APHV
SP	94.1	13.1	2.3	-	-	0.6	Camphor	1.0 20 mm APHV, 20 mm AP, 20 mm HE and 20 mm Inc
							Unac	3.0
Tube	98.1	13.1	-	-	0.02	-	Unac	1.8850 mm Trench Mortar
SP	94.5	13.1	-	-	-	0.8	Camphor	1.3 75 mm APC and 75 mm HE
							Unac	3.4
SP	96.1	13.1	-	-	0.5	-	Unac	3.4 75 mm HE
Square	95.9	15.0	0.3	2.6	-	1.0	Unac	2.2 80 mm Expulsion Powder
Square	98.4	13.1	0.9	-	-	-	Unac	0.7 7.52 mm Rifle Grains (A/T)

Abbreviations: See under Table 44.

Compositions listed in Table 43 are for double-base (NC-NG) propellants analyzed at Picatinny Arsenal during WW II.

(See next page).

Table 43
Double-Base (NC-NG) Propellants

Form	Composition, %						Other Ingredients	Uses
	NC	NN in NC	NG	Case	Acac	Graph		
Tube	58.1	12.3	37.2	5.9	-	-	K sulfates	0.3 37 mm APHV
							Unac	0.5
Tube	69.7	11.9	27.5	1.5	0.2	-	K sulfates	0.6 37 mm APMB
							Unac	0.7
SP	63.7	11.8	28.5	6.3	-	0.1 (incorporated)	Unac	1.5 37 mm HoC
Strip	64.0	12.3	30.0	6.0	-	-	-	37 mm Cerech
Strip	64.0	12.3	30.0	6.0	-	-	-	40 mm Cerech
Strip	63.0	12.2	28.0	9.0	-	-	-	47 mm AP
Strip	63.1	12.4	30.3	6.0	-	-	-	47 mm HE
Strip	62.9	12.2	29.1	7.3	-	-	-	47 mm APCHE
							K sulfates	0.4 and APIN
Tube	61.1	12.0	22.4	12.5	-	-	DNT	0.9 50 mm APC
							Vaseline	1.5
Disc	59.6	12.9	39.0	-	0.7	0.1	K salts	0.6 75 mm HoC
Disc	59.5	13.0	38.7	-	0.8	0.2 (incorporated)	Unac	0.6 75 mm HoC
							Unac	0.8 75 mm HoC (semi-fused)
Square	59.5	12.2	38.6	1.6	-	0.3	-	80 mm HE Mortar
Disc	59.2	13.0	38.5	-	0.6	0.3	Unac	1.4 80 mm HE Mortar
Disc	61.5	12.9	38.1	-	-	-	DPAther	0.4 80 mm CA
Square	58.3	13.1	39.0	0.8	-	0.2	Unac	1.7 80 mm HE
Square	59.6	13.0	38.8	2	0.8	-	DNT	0.4 105 mm How
							Unac	0.4
Square	59.4	12.9	31.4	-	8.9	-	Unac	0.3 105 mm How
Square	55.2	13.0	44.4	-	1.1	0.5	Unac	0.8 150 mm How (Base Charge)
Square	56.8	13.1	40.8	0.3	0.7	0.1	Unac	1.3 155 mm How and 80 mm HE
Square	59.0	13.1	39.0	-	1.0	-	Unac	1.0 155 mm How
Disc	56.5	13.5	41.6	-	0.8	0.2	Unac	0.9 Miscellaneous Mortars
Flake	59.9	13.36	39.0	0.9	-	0.2	-	80 mm Mortar
Square	62.5	12.0	35.0	-	0.2	0.1	DPAther	1.5 150 mm Rocket
							ETPther	1.1
							Unac	1.2

Abbreviations: See under Table 44.

Remarks on Table 43:

The double-base nitrocellulose-nitroglycerin propellants listed in Table 43 were somewhat different from the American and British propellants, as can be seen from the following remarks:

a) In cases in which large amounts of camphor were present, it served not only as a stabilizer, but also as a plasticizer, especially for low-vitrified NC. The amount of NG in such propellants was correspondingly decreased. In other cases where the amount of camphor was small, or even absent, the amount of NG was increased.

b) It has been shown that when castor oil is used in large amounts, it also acts as a thick reducing agent. The same applies to acetic acid (asymmetrical diphenyls). When castor oil was used as a stabilizer, an amount as low as 0.8% was sufficient.

c) Vaseline, present in some propellants, was supposed to act primarily as a cooling agent (to lower the temperature

of combustion and to reduce erosion). It also acted as a stabilizer to a certain extent because the unreacted hydrocarbon present in vaseline combined with the oxides of nitrogen and thus stabilize the powder. It has been found, however, that vaseline is not particularly effective in reducing hygroscopicity.

d) Graphite was used for coating some propellants (see Remarks (c) and (d) in Table 42, b) in propellants of large grain size, such as the 155 mm, 150 mm and 120/45 mm weapons, no graphite coating was used.

e) As in some other German propellants, graphite was used not only as a coating agent but it was also distributed throughout the mass of material. (See Remark (c) in Table 42.)

Table 44 gives compositions of some double-base propellants based on DEGN (diethylenglycidylazinate) and on triple-base (NC-DEGN-NG) propellants.

(See next page).

Double-Base (NC-DEGDN) and Triple-Base (NC-DEGDN-NGu) Propellants

Form	Composition, %					Graph	Other Ingredients	Uses
	NC	KN in NC	DEGDN	Ceas	Acas			
Tube	66.1	11.9	10.2	1.8	0.2	-	Uscac	1.7 37 mm AP Shell
Tube	65.1	12.1	11.5	2.7	-	0.3	K sulfac	0.4 37 mm HE
Tube	66.3	11.8	29.4	2.7	-	0.2	Uscac	1.4 47 mm APHV
Tube	61.4	11.8	29.8	8.4	-	-	Vaseline	0.4 50 mm AP
Tube	61.5	12.0	26.0	7.4	-	0.3	K sulfac	0.5
							Uscac	0.8
Tube	68.7	11.8	28.4	1.5	0.1	-	K sulfac	0.4 30 mm APHV
							Uscac	0.6
Square	58.4	12.6	32.0	-	-	0.3 (incorporated)	NGU	2.9 50 mm HE
							K sulfac (added)	2.7
Tube	60.0	13.1	38.4	-	0.7	0.1	Uscac	0.8 30 mm HE
Tube	57.1	13.9	6.5	0.1	-	0.3	Uscac	1.6 50 mm TM
Tube	60.1	12.4	37.7	0.3	0.7	-	Uscac	1.0 50 mm APHV
Tube	66.4	11.8	29.8	2.3	-	0.2	K sulfac	0.5 30 mm APHV, 47 mm APHV, 57 mm HE and 57 mm AP
							Uscac	0.9
	37.6	12.2	29.8	-	-	0.1	NGU	31.4 42/28 mm Tapered Bore and 42/28 mm APHV
							Uscac	1.1
Tube	65.0	11.9	23.2	8.8	-	0.1	K sulfac	1.5 75 mm AP
							Uscac	1.4
Tube	59.0	12.5	10.5	-	-	0.1	NGU	28.9 75 mm HE
							Uscac	1.5
Tube	62.5	13.0	34.4	-	0.4	0.1	K sulfac	2.5 75 mm HE
							Uscac	0.8
Strip	63.5	12.4	35.9	1.4	0.4	0.1	Uscac	0.7 75 mm HoC, Semi-Fixed
Strip	59.6	12.8	38.6	-	-	0.2	E-Pb/Uscac	1.2 75 mm HoC, Semi-Fixed
							Uscac	0.4
Tube	60.5	11.9	28.2	7.3	-	0.4 (incorporated)	Vaseline	1.7 75 mm Tank Gun
							K sulfac	1.1
							Uscac	0.7
Square	38.4	12.4	31.5	-	-	0.2 (incorporated)	NGU	29.0 75 mm HE HoC, 75 mm HE, Pb 60 and 50 mm HE
							Uscac	0.9
Square, No. 1	62.0	12.4	26.0	7.6	0.2	0.2	E-Pb/Uscac	3.1 76.2 mm AP
							Uscac	0.9
	76.2 mm and some 16 mm weapons were these captured in Russia							
Plate	58.6	12.2	36.9	-	-	0.3	NGU	30.2 76.2 mm HE
Tube	67.2	11.8	28.2	9.3	-	-	Uscac	1.3 88 mm AP
Tube	49.0	11.0	18.5	-	0.2	-	NGU	31.2 88 mm AP
							D-Pb/Uscac	3.2
							E-Pb/Uscac	2.2
Tube	66.7	11.8	28.2	3.3	-	-	Uscac	1.7 89 mm HE
Square	61.6	13.1	37.3	0.3	0.4	0.1	Uscac	0.3 150 mm How (Zones 1-6)
Square	67.1	13.0	34.6	0.4	0.3	0.1	Uscac	0.5 150 mm How (Zone 7)
Disc	58.6	13.0	38.7	0.4	0.5	0.2	Uscac	0.6 150 mm How (Zones 8-9)
Tube	59.6	12.6	33.6	-	-	0.2	D-Pb/Uscac	1.5 75 mm Rocket
							E-Pb/Uscac	3.0
							Uscac	2.1
Tube	61.1	-	35.5	-	2.1	0.2	Uscac	3.3 150 mm Rocket
Tube	59.6	12.5	34.8	-	0.2	0.2	E-Pb/Uscac	1.2 210 mm Rocket
							D-Pb/Uscac	2.0
							Carbazole wax	0.3
							Uscac	1.7
Tube	60.0	-	35.4	-	-	-	Uscac	4.6 300 mm Rocket

(See also G Pulver and Gudolpulver).

Abbreviations: AA Antiaristonic; Alieric; Acm Acetate; Am Ammonium; Ar Armo-piercing; AT Antipiercing; APC Armo-piercing; Capped; AT Anticent; Cen Central; CM Chemical; RMOs DBuDiethylphthalate; DEG Diethylene glycol; DEGD Diethylene glycol Diisocyanate; DNT Diisocyanate; DMA Diphenylmethane; DPHuDiethylene glycol; Et Ethyl; Ethylphenylphenylglycidyl; Phk Gr designation for AA, Graph Graphite; Hc High Explosive; Hc Hollow charge, shaped charge; Hydroxyl; Hydroxyl Hydroxide; Inc Incendary; K (Kannone) Cannon; K salts Potassium salts; LC Long Case; MV Monocyclic; MMT Monocyclic; M Nitrogen; NC Nitrocellulose; NG Nitrocellulose; NGW Nitrocellulose; Pol Gen designation of A/T, PETN Pentaerythritol Tetranitrate; RN Road Note; SC Shot; SP Single Perforation; T Tracer; TEG Triethylene glycol; TETN Triethylene glycol Diisocyanate; TM Trachet; TMT Trinitrocellulose; Ume Unaccommodated.

Remarks on Table 44. (See previous page)

Although the above DEGDN and NGu plus DEGDN propellants were similar in composition to NG propellants listed in Table 43, they had some features which are worth noting, such as:

- a) There was a definite relationship between the percentage of NC and DEGDN used, as the percentage of NC was decreased the amount of DEGDN (which has about the same potential as NG) was increased. It was also noted that decreasing amounts of centralite were accompanied by increasing amounts of DEGDN
- b) The use of low nitrogen content NC, such as 11.8-12%, is DEGDN propellants may be explained by the fact that high N content NC is much more difficult to gelatinize with DEGDN.

c) Several propellants contained about 30% NGu and only about 40% of NC, without any stabilizer. In most of these compositions graphite did not serve for coating but was uniformly distributed throughout the grains. It is to be noted that NGu does not solubilize NC even at low temperatures.

d) All the DEGDN propellants, especially those containing NGs were much cooler burning than the corresponding NG propellants.

e) From the American point of view the DEGDN propellants have the following disadvantages over propellants based on NG:

- 1) They are more volatile
- 2) Less sensitive to flame and thus more difficult to ignite

- 3) More toxic to personnel handling them
- 4) They contain an ingredient (DEGDN) which is more difficult to stabilize than NG.

H. Marnour et al., *Mém. poud.* 35: 280 (1933), gives the composition of a German propellant, used in rounds for 40 mm. calibre canon as follows: NC 42, glucose 14, 210.

63.5% DEGDN 26.5, centralise 8.0 and vaseline 2.0%.

Some information on DEGDON-NC propellants prepared at the Dineberg Fabrik, D A-G may be found in Ref 7. Two of these propellants used in cannons are listed in Table 45a.

The same Ref 7 gives the composition and properties of the DEGDN propellant manufactured by the Wolff Co Plant

DEGDN 17.4, DPhA 0.5, Cent 1.0 and TNT 53.0%. Oxygen balance -16.51% and calorific value 750 kcal/kg.

Some double-base (NC-DEGDN) and triple-base (NC-DEGDN-NGu) propellants manufactured at the Dünberg

Fabrik, Dynamit A-G were described in Ref 5. Their composition is given in Table 43b.

(See next page).

100

Table 45a

NC-DEGDN Prescriptions of Diabetes Enrollments in 1998

Composition and properties	German Designation	
	B5702	Bl4232
NC	29.45	48.59
% N in NC	12.0	12.5
DEGDN	2.645	25.16
Am nitrate	40.00	-
Dicyandiamide	-	25.00
Centralite I	1.00	-
MBN	-	1.00
Hy oxide	-	0.15
Graphite	-	0.10
Moisture	1.10	0.80
Total	101.00	101.00
Oxygen Balance, %	+3.29	-22.44
Calorific Value, kcal/kg	1143	719

Abbreviations: (See under Table 44)

In Ref 6 is described the manu of NC and propellants at the Krümmel Fabrik, Dynamit A-G. while in Ref 8 is described the manufacture of NC and propellants at the following plants: Troisdorf Fabrik DA-G, Ebenhausen Fabrik DA-G, Rottweil Fabrik DA-G and BSLintz Fabrik of Wolff Co.

In the prepn of propellants at the Rottweil Plant the blends consisted of 20 parts NC, N content 12.5%, and 80 parts of NC, N content 13.3%.

Table 46 gives some properties, including the burning characteristics, of several German propellants examined at Picatinny Arsenal during WW II (Refs 4, 10a, 10f and

106).
(See next page).
Remarks on Table 46:

a) In the compositions given in Table 16 only the main ingredients are included. Other components, such as stabilizers, graphite, etc were given in Tables 42, 43 & 44

b) Force of a Propellant ($H \times V$) is a function of its chemical composition

as well as of its composition. The most important variables are total volatile content and web size. The quickness is approximately inversely proportional to the web size. In small, dense granules, the concentration gradient of the

d) The relative quickness of propellants is obtained by determining these burning rates with the aid of a standard

Comparing the burning rates with the rate of a standard. If comparison is made between a German propellant and a standard American propellant, the results are likely to be misleading since the German units made with a heavy

Double-Base (NC-DEGDN) and Triple Base (NC-DEGDN-NG) Propellants of Dünaberg Fabrik, D-6

Form	Composition, %										Calorific value kcal/kg	Uses
	NC	AN in NC	DEGDN	NG	Cres	Acac	Graph	H ₂ O	Other ingredients			
Flake	43.65	13.0	35.80	-	-	0.50	-	0.05	K nitrate	6.50	-	Various Hove
Flake	54.40	13.0	44.50	-	-	0.50	0.05	0.05	K nitrate	6.50	-	Various Hove
Flake	58.03	13.0	31.12	30.00	-	0.50	0.10	0.25	K nitrate	6.50	-	Various Hove
Tube	67.65	12.0	29.00	-	5.00	-	0.10	0.25	-	825	-	88 mm AA and Heavy 100 mm Gun (K18) (Army)
Tube	68.22	12.0	29.23	-	1.70	0.50	0.10	0.25	-	870	-	37 mm AA and 37 mm A/T (Army)
Tube	62.53	12.0	26.72	-	4.00	-	0.10	0.25	Vaseline Phosphate	1.80	700	Heavy Army Field Hows
Tube	61.53	12.0	26.37	-	7.50	-	0.10	0.25	Vaseline Phosphate	1.60	700	100 mm Army Gun (K 18)
Tube	64.08	12.0	27.47	-	5.35	-	0.10	0.25	Vaseline Phosphate	1.85	750	88 mm Army AA Gun
Tube	43.51	12.0	18.64	30.00	-	0.50	0.10	0.25	DPhUret	3.75	750	88 mm Army AA HE Gun
Tube	39.48	12.0	16.92	30.00	-	-	0.10	0.25	DPhUret	4.25	750	88 mm Army AA and AP Gun
Tube	69.32	12.0	14.85	-	3.00	-	0.10	0.15	DNT	10.00	730	88 mm AA and other Army Guns
Tube	60.55	12.0	25.95	-	3.75	-	0.10	0.15	Hydrocell DNT	2.00	750	Various Army Guns
Tube	44.00	12.0	18.85	20.00	-	0.40	0.10	0.15	alpha-MNN	2.50	720	Various Army Guns
Tube	69.38	12.2	25.27	-	5.00	-	0.10	0.25	DPhUret	4.00	820	Naval Guns
Tube	65.59	12.2	25.87	-	9.00	-	0.10	0.25	Phosphate	1.25	730	Naval Guns
Tube	65.71	12.2	25.94	-	2.50	0.50	0.10	0.25	alpha-MNN	7.00	730	Naval Guns
Tube	58.55	12.2	-	-	12.00	-	0.10	0.25	TEGDN	25.10	650	Naval Guns
Tube	35.50	12.2	23.75	40.00	-	0.50	0.10	0.25	K nitrate	4.00	820	37 mm Naval Gun
Tube	42.45	12.0	18.20	25.00	-	-	0.10	0.25	DPhUret	0.70	750	Naval Guns
Tube	60.17	12.6	35.33	-	-	-	-	0.25	Hydrocell	1.50	900	Universal composition for Rocket Launchers
Tube	59.05	12.6	34.82	-	-	0.50	-	0.25	Hydrocell	3.00	865	300 mm Rocket Launcher

Abbreviations: See under Table 44.

Properties of Some German Propellants

Form	Composition, %					Some Properties					Burning Characteristics		
	NC	AN in NC	NG	DEGDN	NG	Uses	Arb (in new)	H	V	Force (lit V)	Δ	A	C
SP Tube	63.46	11.8	28.5	-	-	37 mm hFHC	.0504	881.5	776	674,846	6.62	-	-
Cand	65.7	13.08	20.8	-	-	Antisnark Gun	.0428	890	842	749,580	-	-	-
SP	59.6	12.5	-	38.8	-	Rocket	.246	820.7	705.8	585,602	5.53	-	-
SP	63.0	11.9	-	26.5	-	100 mm K 18	.0537	740.1	407.4	597,556	5.08	-	-
SP	58.5	11.3	-	16.5	34.8	88 mm HEFC	.0577	706.9	680.1	664,762	1.2	-	-
Square	56.0	12.0	-	31.0	32.2	76.2 mm A/T	.0209	877.7	777	681,429	-	-	-
SP	65.0	11.95	-	23.2	-	75 mm HEFC	.0600	712.1	722.5	512,549	4.21	-	-
Sq	63.5	12.4	-	35.9	-	75 mm HEFC	.205	893.8	711.5	695,760	9.0	-	-
Square	59.4	12.9	-	30.9	28.5	75 mm hFHC	.0249	916.6	706.2	643,066	7.26	-	-
Cylinder	59.6	12.55	-	31.6	-	75 mm Leaflet Rocket	.333	856.6	721.0	617,508	-	-	-
Square	40.0	12.4	-	30.3	28.7	75 mm HE Tank	.0261	901	767	691,067	-	-	-
SP	37.4	12.2	-	30.2	31.3	42/28 mm APHV	.0279	883.2	716.2	632,228	5.6	-	-
SP	94.8	12.8	-	-	-	28/20 mm APHV	.0237	829.7	705.8	595,602	0.94	-	-
SP	92.75	13.05	-	-	-	28/20 mm APHV	.0211	829.7	705.8	585,602	0.94	-	-
Square	53.15	13.0	44.4	-	-	150 mm How (Base Charge)	.0067	1235.1	588.6	727,333	9.9	.05	211
Square	61.64	13.1	-	37.3	-	150 mm How (1-6 tons)	.0484	1015.9	685.2	696,094	8.1	.017	167
Square	62.31	13.0	-	36.6	-	150 mm How (7 tons)	.0515	995.6	696.7	692,242	8.4	.009	144
SP Disc	59.6	13.0	-	38.7	-	150 mm How (7-8 tons)	.0722	989.4	704.5	697,057	8.8	.21	158

Abbreviations: A Constant called Vivacity; C Rate of evolution of hot gases at a pressure of 20,000 psi in liters at atmospheric pressure / sq cm of surface / second; H Heat of Combustion in kcal/kg; P Pressure of propellant gases in psi; V Volume of gases liberated in l/kg; Δ Burning rate (quickness) of the propellant at 20,000 psi in inches/sec; (NAV) Force or Time-mechanical Potential.

Other abbreviations are given under Table 44.

about distance along the bore of the gun. On the other hand, in American guns with a lighter breech the shells are designed to develop the maximum pressures more slowly and after the shell has travelled a greater distance along the bore of the gun.

a) In the relation of quickness to composition, it may be noted that the single-base propellants are the slowest and are comparable to those double-base propellants which contain NG. Propellants containing NG are usually the fastest, followed by DEGDN propellants. In some cases, however, DEGDN propellants are faster than those containing NG. This is usually the case when the NC in a DEGDN propellant is of considerably higher nitrogen content than that used in a corresponding NG propellant.

f) The burning rate of the German 210 mm rocket propellant was given equal to $0.35 \text{ (CP)} \times 10^{-2}$ while the corresponding value for the standard American double-base 7/8" stick propellant is 48.6×10^{-2} . This means that the rate for the American propellant is about 65% greater than for the German propellant.

g) Experimental procedures for the determination of the burning rate of propellants are described in Pic Area Tech Rept 1235 (1943).

h) Methods of composition of propellants of propellants are given in the Du Pont, Bunsdale Laboratory Memorandum Report 31.

Reference (Propellants):

1) Adairhall, Explosives, Churchill, London, v1 (1977)

v2 (1917) and v3 (1937)

2) H. Bunsdale, Das rauchlose Pulver, W. de Gruyter Berlin (1926).

3) A. Stettbacher, Schiess- und Sprengstoffe, J.A. Barth, Leipzig (1933)

4) Collective, PB Rept 11,544 (1943)

5) G.N. Schilling et al., PB Rept 925 (1943)

6) L. Nutting et al., PB Rept 16,666 (1945)

7) F.J. Krieger, M. Pleaser, PB Rept 7826 (1943)

8) R. Ashcroft et al., BIOS Final Report 833 (1944), Item 2

9) H.H. Mphike, CIGS Report 31-68 (1946), Report on Visit to Dünaberg Factory of D. & A.

10) A. Stettbacher, Spreng- und Schiessstoffe, Rascher, Zürich (1948)

11) Plantany Arsenal Technical Reports:

a) Collective, 1282 (1943) (Fertilizer Propellants)

b) A.B. Schilling, 1358 (1944) (Propelling Charge for 155 mm Separate Loading Ammunition)

c) A.B. Schilling, 1439 (1944) (Separate-Loading Propelling Charge Assembly for 105 mm Recoilless Gun, LC 41)

d) J.P. Wardlaw, 1443 (1944) (Propelling Charge for Separate-Loading 100 mm Gun, K 18)

e) A.B. Schilling, 1453 (1944) (Propelling Charge for 210 mm Separate-Loading Ammunition)

f) Collective, 1456 (1944) (Foreign Propellants)

g) W.R. Tomlinson, Jr., 1555 (1945) (Chemical Composition of Materials used in German Ammunition)

Propellants: Artillery. According to H.H.M. Pike, CIOB Report 51-68 (1946), pp 4-8 and tables, the following types of propellants were used by the Germans in their artillery weapons:

- [illegible]

Table 47 gives composition and some properties of common artillery propellants used during WW II by Germans. (See following pages).

Propellants, internal Ballistic Data is given in tables at the end of CLOS Report 31-68 (1946).

Propellants, Rocket. See Rocket Propellants.

Propellants, Stability of. The stability characteristics of some German propellants were determined during WW II at Picatinny Arsenal and described in Technical Report 1436 (1944).

In cases where sufficient material was available, both the 120° and 134.5° Heat Tests were made. The results of tests showed a tendency toward greater stability for those propellants which contained a stabilizer-gelatinizer (such as centralite) in combination with another stabilizer, such as acardite.

Sufficient amounts of propellants were not available for tracking a definite conclusion concerning the merits of disubstituted urethanes in combination with acrolein.

Propellants containing NG, DEGDN and NG-DEGDN proved to be of satisfactory stability, judging by the 120 Heat Test of the U.S. Army (the test paper should not turn a salmon pink color in less than 40 minutes).

As to the single-base propellants, only a few of the German propellants met the U. S. Army Specification which requires that the test paper in the 134.5° Heat Test and the not rate salmon pink in color in less than 45 minutes.

Propellant Charge in Fixed and Semi-Fixed Ammunition. According to E. Engleburg (The Ordnance Sergeant, May 1944, p 321), German propelling charges may be subdivided into two main classes:

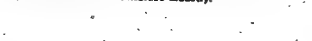
- [illegible]

Propellant Grains and Their Dimensions. The following typical German propellants are listed by H.H.M.Pike in C/OS Report 11-68 (1946), pp 4-5 and emblem:

- a) Tubular (Rorippa sp.), designated as RLP (3 × 1 × 0.5), consisted of tubes 810 μm long having external and internal diameters of 13 mm and 4.3 mm respectively (Fig. 1a); (Rorippa sp.), designated as RLP (10 × 10 × 0.6), consisted of grains 100 μm long, 10 mm wide and 0.6 mm thick;
- c) Flake (Blattenschuppe), designated as BLP (3 × 1 × 0.5), consisted of grains 3 mm long, 3 mm wide and 0.6 mm thick;
- d) Disc (Plättchenpulver), designated as PIP (50 × 50 × 0.2), consisted of discs 50 mm in diameter and 0.2 mm thick;
- e) Ring or annular (Ringpulver), designated as RLP (50 × 50/10), consisted of grains 0.5 mm thick, 50 mm in diameter and a central hole of 10 mm in diameter.

TABLE C
Artillery Personnel[illegible]

...a bomber.

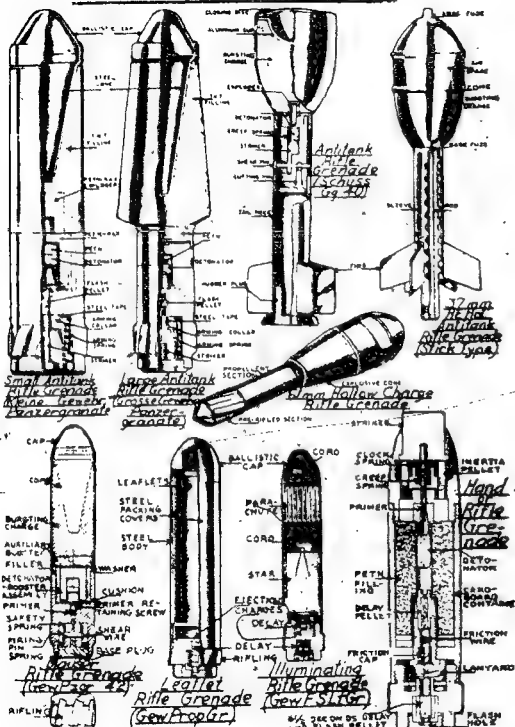


ESTABLISHED PRODUCT WAS GRATED FINELY IN ONE FORM OF SMALL

References: Intelligence - Bulletins, U S War Department, Washington, D C , Vol III, No 3 (1945), pp 74-79 and Vol III, No 7 (1945), pp 9-16.

Ger 159

RIFLE GRENADES



Ger 160

- 2) Marshall, v.1 (1917), p. 491
3) Colver (1918), p. 141.

Table 52
Roburites

Components and Some Properties	Designation		
	I	II	III
Am nitrate	87.5	71.5	55.0
K nitrate	-	5.0	9.5
K permanganate	0.5	0.5	0.5
Am sulfate	5.0	-	-
m - DNB	7.0	-	-
TNT	-	12.0	12.0
Flour	-	6.0	6.0
Nb chloride	-	5.0	7.0
Am chloride	-	-	5.0
Magnesium	-	-	5.0
Trausal Test, cc	-	325	257

Röchling Anticoncrete Projectile (Röchlingsgranate 42 Beton, patented as RÖG 42 Bet. According to German photographs available at the Picatinny Arsenal and Aberdeen Proving Ground Museums, it was a subcaliber shell which resembled in appearance a "arrow projectile", except that instead of the fin assembly of the arrow shell it had a discarding flange serving as a driving band. The front flange acted as bourrelets. These projectiles were fired from regular guns, such as caliber 21 cm and 14 cm. The 21 cm shell weighed 193 kg and was 2.1 m long. The corresponding characteristics for the 14 cm shell were 913 kg and 3.7 m.



RÖCHLING PROJECTILES

The shells were designed and manufactured by the firm of Röchling at Saarbrücken, Saar.

- References:
1) K.F. Kämpf, Museum of Aberdeen Proving Ground, Md; private communication.
2) H.H. Bullard and G. Coghlan, Picatinny Arsenal Museum; private communication.
(See also Arrow Projectiles and German Projectiles).

Rocket (Rakete). German rockets of WW II were propelled either by solid propellants (such as cellophane nitrocellulose double-base NC-NG propellants) or by liquid propellants. The liquid propellants consisted of combinations (such as alcohol, benzene, aniline, gasoline etc.) and oxygen carriers, such as liquid oxygen, hydrogen peroxide, nitrogen peroxide, nitric acid, etc. (See under Rocket Propellants).

The following rockets were briefly described in Ref. 3. (Some information on these rockets may be found in Refs. 1 and 2).

- a) Butterfly (Schmetterling) Rocket: (Hill) (Hs. 297) (Ref. 3, p. 156) (See under Guided Missiles)

The total weight was 19 oz, overall length 238 mm, length of stem 102 mm and its diameter 30 mm, length of head 116 mm and its max diameter 40 mm. Its burning and propellant charges, as well as its pressure, detonator and booster were the same as for the 40 mm grenade (Ref. 1, p. 9, Ref. 2, p. 11 and Ref. 3).
c) of mm Antitank Rifle Grenade, briefly described on p. 331, Ref. 3, was similar in construction to the previous grenade. Its overall length was 244 mm.
d) Leaflet Rifle Grenade (Gewehr Propagandagrüne) was fired from the rifled 10 mm diameter cup (Schalenbecker) which could be fitted to most types of German rifles. It consisted of a cylindrical steel body (with a flattened base) containing a delay fuze, a disc cylindrical container for the pamphlets and an ejecting charge for this cylinder. On firing the grenade, the propellant gases ignited the delay fuze and, after about 9 seconds of delay, the fuze fired the ejecting charge. The resulting detonation blew off the cap and forced the leaflets out the nose. Total weight of grenade 8 oz, overall length 5.7" and range 500 yd (Ref. 3, p. 330).

e) Illuminating Parachute Rifle Grenade (Gewehr Fallschirmscheinwerfer) consisted of a thin-walled cylindrical body, within which was another container which housed the parachute and illuminating star. The test of grenade contained two delay pellets and two ejection charges. When fired the fuze from the propellant gases ignited delay (1) and after 5.5 sec of light ejection charge (1) was initiated. The pressure of the gases forced out the nose, the container (which held the parachute) and the star. At the same time, delay (2) was ignited and after it burned through (2 seconds) the ejection charge (2) became initiated. The resulting gases ejected the parachute and the star from the container and ignited the star. It was claimed that distances up to 650 meters could be illuminated by this grenade (Ref. 3, p. 331).
(See also Flareparatroo and Paratro Grenades).

- References:
1) A.J. Dere, The Ordnance Sergeant, October 1945, pp. 8-10;
2) Anna, TM 9-1985-2 (1953), pp. 331-39
3) Picatinny Arsenal Technical Report:
a) A.B. Schilling, No 1342 (1944)
b) A.B. Schilling, No 1396 (1944)
c) A.B. Schilling, No 1345 (1945)
d) F.G. Haverak, No 1307 (1945)
e) F.G. Haverak, No 1309 (1945).

Rifle (Gewehr). See under Weapons.

Rifing of Weapons. See general section.

RLCS (Raketenleuchtgerät Scheinwerfer) Rocket Illuminant Simulating Device. See under Pyrotechnic (Incendiary) Devices and also in CIGS Rep 35-56 (1943), p. 21.

R. mine 42. See under Landmines and also in TM 9-1985-2 (1953), p. 472.

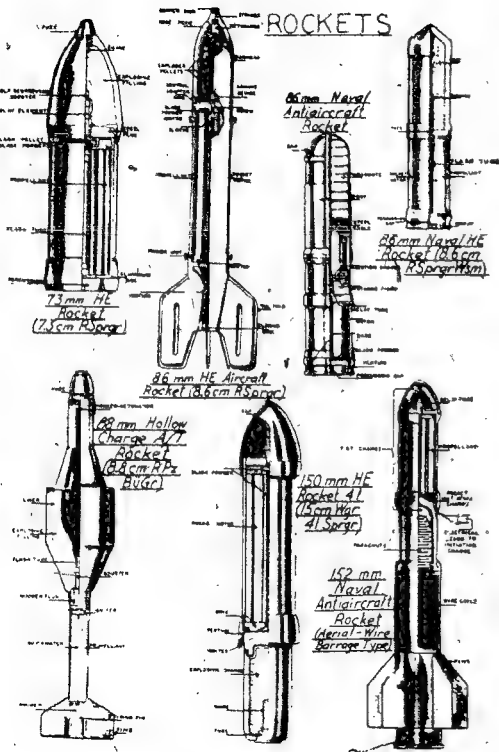
Roburite (Roburite)-A type of permissible explosive patented by Rodt about 1886. The earliest type consisted of Am nitrate 90 and dinitrophenol 10%. It was claimed by the inventor that a nitroated chloro-compound gave a higher velocity of detonation and greater power than the corresponding nitro-hydrocarbon. The above Roburite was sensitive to friction, when ignited with a flame or a spark it burned without explosion.

Table 52 gives the composition and some properties of several Roburites.
(See next column).

- References:
1) Daniel, Dictionnaire des Matières Explosives, Paris (1902), p. 687

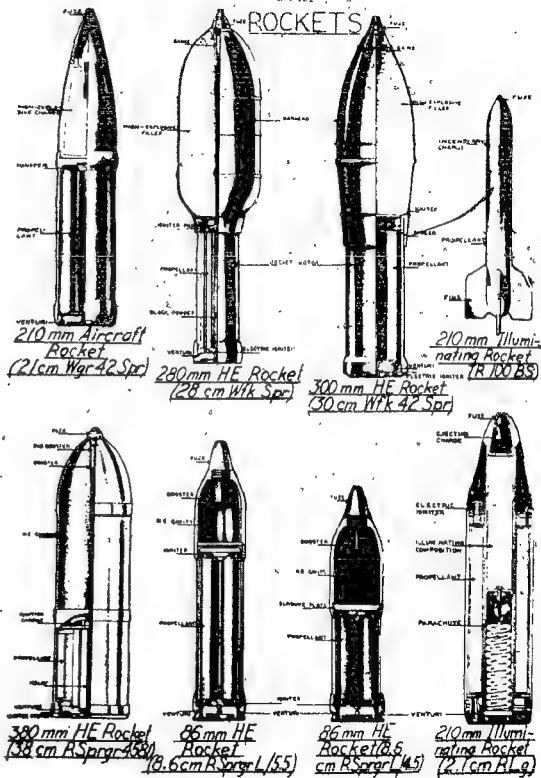
Get 101

ROCKETS



Get 102

ROCKETS



- b) Daughter of the Rhine (Rheinischer) Rocket (Ref 3, p 226)
- c) Fire Lily (Feuerlily) Rockets 7-25 and F-55 (Ref 3, p 224)
- d) Great Russian Rocket (Ref 3, p 225)
- e) Henschel Rockets H-755 and H-1250 (Ref 3, p 200)
- f) Long Range Rockets A-9 and A-10 (Ref 3, p 235)
- g) Radio-Controlled Glider Bomb PC 1400 FX (Ref 3, p 195)
- h) Rockets V-1 and V-2 (Ref 3, p 203)
- i) Rocket X-4 (Ref 3, p 214)
- j) Teufel Rocket (Biliquid) (Ref 3, p 225)
- k) Wasserfall/Wasserfall Rocket C-7 (Ref 3, p 219)
- l) 175 mm Propaganda Rocket (23 cm Propagandagranaat) (Ref 3, p 234) and 75 mm HE Rocket Shell (7.5 cm Raketenartillerie) (Ref 3, p 235)
- m) 80 mm HE Rocket Shell (8 cm Raketenartilleriegranaat) (Ref 3, p 237)
- n) 85 mm HE Rocket Shell (8.5 cm Raketenartilleriegranaat) (Ref 3, p 239), 80 mm R Sgr L/4.5 Rocket (Ref 3, p 256), 86 mm Illuminating Rocket (Novall) (Ref 3, p 240) and 80 mm Antiaircraft Rocket (Novall) (Ref 3, p 241)
- o) 88 mm (w/c, A/T) Rocket (shaped charge antitank) (Ref 3, p 242)
- p) 150 mm HE Rocket (spine-mounted) (Ref 3, p 245) and 150 mm Smoke and Chemical Rocket (spine-mounted) (Ref 3, p 245)
- q) 112 mm Antiaircraft Rocket (spine-mounted) (Ref 3, p 247)
- r) 200 mm Antiaircraft Rocket (spine-mounted) (Ref 3, p 240)
- s) 210 mm HE Airborne Rocket (spine-mounted) (Ref 3, p 242) and 210 mm Illuminating Rocket R-L (Ref 3, p 250)
- t) 250 mm HE Rocket (spine-mounted) (Ref 3, p 250)
- u) 300 mm HE Rocket (spine-mounted) (Ref 3, p 251)
- v) 320 mm Incendiary Rocket (spine-mounted) (Ref 3, p 252)
- w) 580 mm HE Rocket (spine-mounted) (Ref 3, p 254)
- x) 800 mm HE Rocket (spine-mounted) (Ref 3, p 255)

Abbreviations: HE High explosive; HE Half-ton charge (See also Guided Missiles).

- References:
- 1) A. Detsch, Leo Anton Seibert, Altmann, Berger-Lorenz, Paris (1947), pp 140-144
 - 2) A. Seibert, "Kriegs- und Schützenartillerie, Raketen, Zürich (1948), pp 30-37
 - 3) Dept of the Army Tech Manual TM 9-1085-2, (1955) pp 195-246
 - 4) J. C. Tschinkel, Chem Eng News 32, 2542-2547 (1954)

The following Pictorial Annual Technical Reports were derived in German rockets:

- 1) A. B. Schilling, Pic Area Tech Rep 1427 (1944), 30 mm Rocket type rocket
- 2) A. B. Schilling, Ibid 1568 (1945), Wehrmacht and Fuses of A-4 Rocket (Called also V-2 Rocket)
- 3) V. A. Underhill, Ibid 1617 (1951), Evaluation of Some Rocket Propellants Used in V-2's (Confidential)

Note: None of the confidential reports were used as sources of information for this work.

The following CLOS Reports contain some information on German rockets:

- 1) G. Collins, CLOS 29-50 (1949), Rockets and Guided Missiles. (Based on the article of Dr. W. von Braun, Survey of Development of Liquid Propellant Rockets in Germany)
- 2) F. G. Ewing & M. M. Miller, CLOS 29-45 (1943), Luftwaffe-Raketenpanzer-Hermann Göring (Rocket)

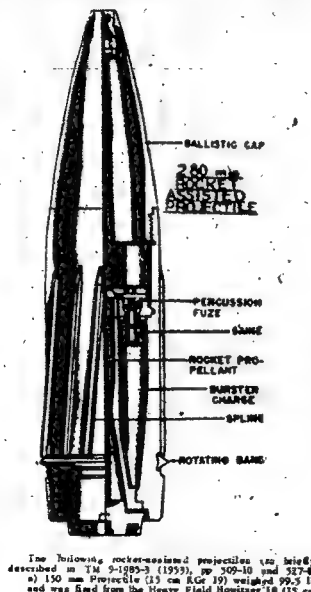
102 R.C. Staff, CLOS 30-115 (1945), Rocket Power Plant Design and Constructed by Walter Wirtz, Kiel

113 F. J. Ewing & M. M. Miller, CLOS 31-13 (1945), Kammer and Rocket Motors (Rockets)

123 H. J. Eppa, CLOS 32-56 (1945), Pyrotechnic Antiaircraft Rocket (Includes description of pyrotechnic rocket: 15 cm RSG-5, 15 cm RLG-5 and 15 cm Smoke Rocket)

13) A. B. Schilling, CLOS 32-114 (1945), 21 cm RLG Rocket (Plan)

Rocket-Assisted Shell: A projectile which contained a rocket propellant in a special device attached to the base of the shell was developed and used during WW II. The shell was fired in a regular manner from a 8 inch gun but during the flight the rocket composition became ignited and the shell started to function as a rocket. This method of propulsion increased the range of the shell from 38 to 60 miles without appreciable increase of dispersion. Reference: PB Rep 525 (1945), p 19.



a) 18. Its cartridge case (semi-fired) contained 13.54 lb of suboxide, diethylene glycol damage type propellant

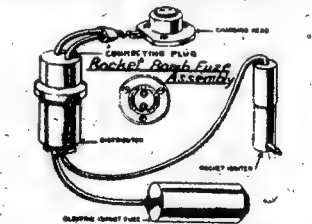
b) 280 mm Projectile (28 cm RLG 4331) weighed (without rocket ignition fuse) 5465 lb and was fired from the Railway Gun (28 cm RLG 4331). Its propellant charge was 45 lb of double-base propellant, and the burning charge was 10% of unknown HE. The shell was provided with a rocket ignition fuse (ZiZ S/40) which functioned after 19 seconds to ignite the rocket and with two fuses (A2, A33), and two PEIN boosters (ZZ44 G/C/889) which initiated the burning charge on impact.

c) In CLOS Rep 30-115 (1945), pp 26-27 and enclosure 20 are briefly described the Rocket Assisted Tail-Off-Usser, designated as RLG 205 and RLG 206.

The following unclassified Pictorial Annual Technical Reports describe some rocket-assisted shells which were examined during WW II.

- 1) A. B. Schilling, 1604 (1946), 105 mm Rocket-Assisted, HE
- 2) A. B. Schilling, 1601 (1946), 105 mm Rocket-Assisted, HE
- 3) A. B. Schilling, 1606 (1946), 128 mm Rocket-Assisted, HE
- 4) A. B. Schilling, 1607 (1946), 150 mm Rocket-Assisted, HE
- 5) A. B. Schilling, 1608 (1946), 150 mm Rocket-Assisted, HE
- 6) A. B. Schilling, 1609 (1946), 150 mm Rocket-Assisted, HE
- 7) A. B. Schilling, 1610 (1946), 150 mm Rocket-Assisted, AP.

Rocket Bomb Fuse Assembly, described on pp 16-21 of TM 9-1085-2 (1955) operated as follows: On release from the aircraft the electric charge passed from the charging lead to the distributor and thence directly to the bomb fuse. Then, after a delay the current passed to the



rocket propellant ignited. During the flight, the rocket was ignited and when the bomb hit the target the impact initiated the fuse. After a short delay (for penetration purposes) the burning charge was detonated.

Rocket Bullet. According to CLOS Rep 35-20 (1946), pp 6, 64 & 7, a 9 mm rocket missile was made during WW II by the Deutsche Waffen- und Munitionsfabrik A-G, L. Bock, a device is enclosed in CLOS Rep 35-20 but no description given.

Rocket Launcher or Projector (Raketenwerfermaschine oder Wurfbatterie). According to the Intelligence Bulletin, War Department, Washington, D.C., vol 5, No 7, March 1945, p 39, the five German rocket launchers were known as Wurfbatterien. Heavy throwing apparatus 400 and 1400 mm were used. Each of them could fire either 200 mm or 320 mm rockets, weighing 180 and 150 lb respectively. The 320 mm HE rocket also could be fired from these launchers. The 300 40 launcher consisted of a wooden frame (Hauptgestell 40) on which were placed wooden shipping crates containing rockets. The frame was inclined at

the desired angle and the rockets were fired directly from the crates.

The 300 41 launcher consisted of a frame of steel tubing (Wurfbatterie 41) on which could be placed the rockets. The rockets were fired from the Wurfbatterie 41. The Wurfbatterie 41 consisted of a sectioned frame mounted on a rocket carrying cradle, track (three on each side), the latter then inclined at the required angle of firing. One of the most important rockets to be used was the 15 cm Nebelwerfer 41 (literally "smoke crawler"), nicknamed by the U.S. soldiers "Screaming Mermers". It consisted of two-wheeled carriage with a split rail. The crew of a light mortar platoon (one platoon) could fire 8 rockets in 2 minutes then discharged the rockets in a salvo (one salvo 8 minutes) by remote control. The maximum range of these rockets was 8,000 yd.

Similar to the 15 cm Nebelwerfer 41 was the five-wheel 21 cm Nebelwerfer 42 which fired 8 inch rockets as far as 8,600 yd. Note: None of the Nebelwerfers were accurate and for this reason were not very suitable for launching HE of smoke rockets. Besides using these launchers for rockets in lay down smoke concentrations, they were also suitable as projectiles for gas-loaded (chemical) rockets. In both cases no accuracy in fire was required.

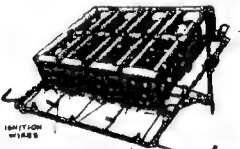
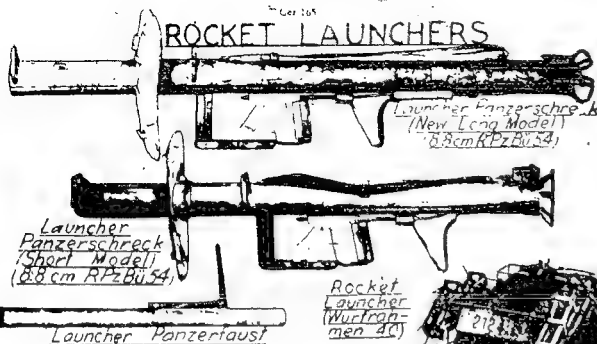
In order to give their larger rocket projectiles greater mobility and speed of fire, and to increase the accuracy of fire of the rockets the Germans mounted the steel frames of the Nebelwerfer 41 and 42 on pneumatic wheels. The resulting weapons were called 280 32 cm Nebelwerfer 41 and 300 cm Nebelwerfer 42. The steel shipping crates containing rockets were loaded in the frames and then, when ready to fire, the crew (seven men per pair of launchers) took cover in non kill trenches in the rear of the launchers and fired the rockets. The rockets were fired at the right side of the weapon and the crew took cover in a separate salvo by remote control. It took about 5 minutes to reload the weapon. The maximum range for the 280 cm HE rocket was only 2,100 yd and for the 320 mm incendiary rocket 2,400 yd. The range for the 300 cm rocket is not given.

Discontinued with the slow rate of fire of the above launchers, the Germans in 1942 began the development of a weapon called the 15 cm Panzerwerfer 42 (150 mm anti-tank weapon 42). It consisted of two tanks of 15 cm caliber (one for launch and one for firing) mounted on a motorized half-track. Since the crew did not need to go into the trenches, but could take cover in the vehicle instead, the rockets could be fired somewhat faster than from the Nebelwerfer 41.

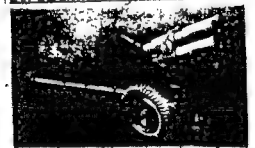
According to TM 9-1085-2 (1955), p 17, ball-mounted projectiles carrying up to 41 rocket rounds were developed by the Germans to effect a greater range of fire. Reloading of these projectiles was carried out mechanically.

The same TM 9-1085-2 mentions or briefly describes the following rocket launchers used during WW II:

- a) A two-wheel cradle type launcher for the 117 (He 297) Schmetterling rocket-propelled missile (p 201)
- b) A 100 mm type launcher (100 mm rocket) (p 201)
- c) A 150 mm type launcher (150 mm rocket) (p 201)
- d) A 150 mm type launcher (150 mm rocket) (p 201)
- e) A 150 mm type launcher (150 mm rocket) (p 201)
- f) A 150 mm type launcher (150 mm rocket) (p 201)
- g) A 150 mm type launcher (150 mm rocket) (p 201)
- h) A 150 mm type launcher (150 mm rocket) (p 201)
- i) A 150 mm type launcher (150 mm rocket) (p 201)
- j) A 150 mm type launcher (150 mm rocket) (p 201)
- k) A 150 mm type launcher (150 mm rocket) (p 201)
- l) A 150 mm type launcher (150 mm rocket) (p 201)
- m) A 150 mm type launcher (150 mm rocket) (p 201)
- n) A 150 mm type launcher (150 mm rocket) (p 201)
- o) A 150 mm type launcher (150 mm rocket) (p 201)
- p) A 150 mm type launcher (150 mm rocket) (p 201)
- q) A 150 mm type launcher (150 mm rocket) (p 201)
- r) A 150 mm type launcher (150 mm rocket) (p 201)
- s) A 150 mm type launcher (150 mm rocket) (p 201)
- t) A 150 mm type launcher (150 mm rocket) (p 201)
- u) A 150 mm type launcher (150 mm rocket) (p 201)
- v) A 150 mm type launcher (150 mm rocket) (p 201)
- w) A 150 mm type launcher (150 mm rocket) (p 201)
- x) A 150 mm type launcher (150 mm rocket) (p 201)
- y) A 150 mm type launcher (150 mm rocket) (p 201)
- z) A 150 mm type launcher (150 mm rocket) (p 201)



Rocket Launcher (20.7 cm Wurfgerät 41)



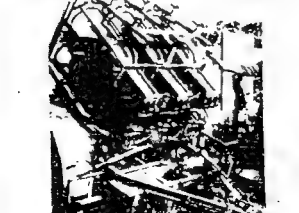
Launcher (15 cm Nebelwerfer 41)



Launcher (15 cm Panzerwerfer #2)



Launcher (23.5 cm Nebelwerfer 41)



Launcher (Do Werfer or Wurfgerät)

b) A four frame launching stand (Wurfgerät) for the 750 mm HE rocket (28 cm Wd Sp) (p 151). (See also under Weapons).

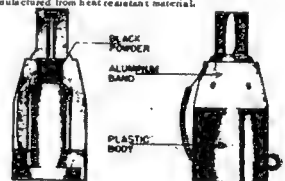
Rocket Projectile. See Rocket-Assisted Shell.

Note: Rocket-assisted projectiles were fired either from howitzers or guns. For instance the 15 cm RGr 19 was fired from the 15 cm w/ff 18 (heavy field howitzer 15) and the 28 cm RGr 413 was fired from the 28 cm RGr 413 (medium gun 70). See TM 9-198-3 (1955), pp 300 A, 329 1.

Rocket Propellant. According to Litomski, Przemysl Chemiczny 77 (4), p 487 (1948), (translated by Dr. Litomski), the Germans used solid double-base propellants containing nitrocellulose and nitroglycerine in their smaller rockets. The larger types, such as the V-2, used liquid propellants consisting of a fuel (such as alcohol, hydrazine, fuel oil, etc.) and an oxygen carrier (such as hydrogen peroxide, nitric acid, tetroxymethane, etc.). Mixtures of easily oxidizable organic liquids with hydrogen peroxide of 80-95% concentration were the most widely used. Hydrazine peroxide could also be used as the driving force, without any fuel, because the heat liberated according to the reaction of decomposition.

$H_2O_2 \rightarrow H_2O + \frac{1}{2}O_2$ 2150 kcal was sufficiently great. Water (vapour) and oxygen served as driving forces.

Rocket Propellant Igniter ERZ 39. briefly described on p 623 of TM 9-198-3 (1951), fitted into one of the venturi of the 15 cm and 21 cm rockets. Its body, made of a plastic with an aluminum band around the shoulder, contained an igniter bridge from which ran two wires. One wire was connected to the aluminum band around the shoulder and the other to a metal disk in the base of the fuse. Just above the igniter bridge was located a black powder charge. When an electric current passed through the bridge, it ignited the black powder, which in turn ignited the propellant. This modified version of igniter (ERZ 39) was manufactured from heat resistant material.



Rocket Propellant Lubricating Coating. In order to prepare a stick of propellant, so that it would burn from an end and not on the sides, the lining was made that it was sufficient to cover the sides of such stick by dipping it twice into a special composition developed at the Dusseldorf Plant of the Dynamit A.G. This composition consisted of: polyethylene 5, stibylene 1/25, 5, Bu50 10, methylacrylate 5 and water 40%.

Get 100.

References: A.A. Swanson and D.D. Sager, JUS Rep 70. 4 (1948), p 4 (As reported by Dr. H. Litomski).

Rocket Propellants, Liquid. The following liquid rocket propellants were used by the Germans during WW II.

1) Concentrated hydrogen peroxide and C-Stoff was used in the da 1401 batter rocket (surface to air). Note: C-Stoff is a 50/50 mixture of methanol and hydrazine hydrate, N₂H₄ (p 3).

2) Concentrated nitric acid and Viscol was used in the Fieseler Fi 103 Rocket, Rheinstetten A-1 Rocket and Westwall Rocket.

Note: Viscol is (in) leavy ether:

1) Concentrated nitric acid and Tocka were used in the Luftwaffe C-4 Rocket.

Note: Tocka is a mixture of aniline, monomethylamine, dimethylamine, p. aniline, naphthalene, methylaniline and isobutylamine.

4) Concentrated hydrogen peroxide with Kpermanganate was used in the Heinkel Rocket.

5) Liquid oxygen, alcohol and water were used in the V-2 Rocket and in the Heinkel-55 Rocket.

Note: The noncombustible substance, water, was incorporated in order to see the flame temperature as low as possible as to reduce the mechanical strain on the motor without sacrificing too much performance. It was found that the addition of 25% of water to absolute alcohol lowered the chamber temperature 7%, while the exhaust velocity was lowered on 1.15%.

6) Concentrated nitric acid, aniline and triethylamine were used in the Schmeisser Ha17 Rocket.

7) Concentrated nitric acid and butyl ether were used in the Taifun Rocket.

8) Compressed oxygen and gasoline were used in the V-1 Rocket.

Note: In addition to these the following substances were used in liquid fuels: aniline, ethyldiamine, ethyldiamine, acetaldehyde, naphtha, gasoline, dimethylamine, monomethylamine, triethylamine, isobutylamine, etc. In some of these liquids, such as aniline, Viscol, ethyl pyrocatechol (Brenskatechin in German) was dissolved. References:

1) H. Garmann, Veltzraumfahrt 6, 134-9 (1951), Jato and Auxiliary Rocket Propellant Plants.

2) K. G. Guld, Development of the Guided Missile, Philosophical Library, N.Y. (1952), p. 112-127.

3) J. G. Tschinkel, Chem. Eng. News 32, 2362-87 (1954).

Propellants for Rockets and Space Ships.

Rocket Propellants, Solid. All known German propellants of WW II were based on NC and a nitric ester, such as NO, DEON, or TE50N.

Table 53 lists some of the rocket propellants examined at Picatinny Arsenal during WW II.

(See next page).

Donnan and Donovan (Ref 5) give the burning rates (in inches per second) at various pressures for the solid propellant used in the 210 mm Rocket (See Table 54 on next page). The composition of the propellant is given in Table 51.

The same investigators give the rates of burning for the jet-assisted Take-Off-Lift Propellant listed in Table 51 (See Table 55 on next page).

Table 53
Rocket Propellants, Solid

Rocket Propellants, 1941										
Form	Composition, %					Graphite	Other Ingredients	Uses		
	NC	NC in HN	NG	DEGN	CBR				ACR	
SP	62.5	12.0	33.0	-	-	0.2	0.1 (sacer)	EPAliver Use	1.5 0.9%	150 mm HE Rocket
SP	58.7	12.7	-	35.3	-	6.2	0.3 (sacer)	EPAliver DPAUse Use	1.3 2.5% 1.7%	210 mm Rocket
	64.1	12.7	12.7	-	0.8	2.4				210 mm Rocket (Igniter Pad)
	89.2	12.7	-	5.3	0.9			DPA Use	2.6 2.0%	210 mm Rocket (Head Igniter Diaphragm)
Cyl	59.6	12.5	-	33.6	-		0.2 (sacer)	DPAUse DPAUse Use	1.5 3.0 2.1%	75 mm Launcher Rocket
	64.7	12.0	-	29.3	-	6.2	0.1	EPAliver DPAUse (TiO ₂ BaSO ₄)	3.5 1.3 0.9%	Jet Assisted Take Off Unit

[illegible]

Note: Some rocket propellants and igniters analyzed at Picatinny Arsenal are listed under Propellants (See Tables 43, 44, 45b and 48).

Table 54:
Burning Rates of 210 mm Rocket Propellants
(Inches per second)

Temp °C	Pressure in psi				
	500	1500	2500	3500	4500
-25	-	0.30	0.42	0.55	-
+50	0.21	0.43	0.55	0.73	0.93

Table 33
Burning Rates of the Jet-Assisted-Take-Off-Unit Propellants

Temp °C	Pressure in psi					
	500	1000	1500	2000	3000	3500
-23	0.13	0.18	0.25	0.33	0.45	0.48
+50	0.22	0.27	0.39	0.47	0.59	0.66

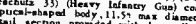
According to Ref 4, the Reissdorf Fabrik W 9 A-G manufactured during WW II several types of rocket propellants. Their compositions are given in Table 36

Table 56 - 1
Rocket Propellants, Solid of W A S A -G

Compounds and some properties	Distillation				
	R.G.	R.W.	Z15	Z16	Z16
Diisocyanate (DICI)	19.80	17.70	49.16	63.25	34.94
X-Minor in HC	12.3	12.5	12.7	82.5	12.5
Dicyanoglycol diisocyanate	35.30	18.00	50.00		14.30
Diisocyanate glycol diisocyanate (TEGDI)				22.00	
Polycarbonate isocyanate (PETN)			30.00	6.00	6.00
Ethylphenylurethane	1.10				
Diphenylmethane	0.90				
Diethylglycolurethane		3.00			
Acetone 2, 2'-COOH ₂ HC ₂ H ₅	0.50	0.50	0.75	0.50	0.50
Glycolurethane		0.30	0.10		
Magnesium salt	0.10	0.50	0.65		
HC Phenyl Wax E	0.63				
Potassium nitrate	0.33				
Lignin					0.7
Hypocellulose	1.50			0.75	
Transisocyanate (TNY)				1.00	12.5
Diacetone urethane (DNT)				0.50	0.90
Miscellaneous (not included in total)	1.00	0.65	1.00	1.00	0.9
Oxygen balance	-7.15	-7.94	-6.01	-6.31	-6.99
Carbonic Yield (at 100)	70.8	89.7	107.6	108	126

' Titration 2/14/78 (TIO)

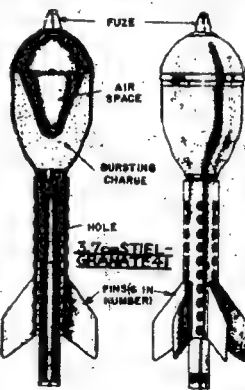
References:

- 14) A.J. Phillips, *Pic Area Tech Repts* 1282 (1945); *Ibid* 1456 (1949).
- 2) Collective Data on Foreign Ammunition, *PB Repts* 11,544 (1945).
- 3) M.D. Davis & F.J. Donovan, *Copied Enemy Propaganda, OSRD of HRCB*, 2d Ser., Sec. 1, *Chem. and Rept*, *Seizure P*, No. 10.2 (1945) (Unclassified) (OSRD 585).
- 4) F.J. Krueger & M. Plossner, *PB Rept* 1826 (1945), p. 6.
- 5) F. Bellinger, *Ind Engng Chem* 38, 3, pp. 160-9 (1946).
- 6) R. Levy, *Chimie et Industrie*, 57, 121 (1947).
- 7) J.G. Tschinkel, *Chem Eng News* 28, 2382-87 (1954).
- 8) *Propaganda for Rockets and Space Ships*
- 9) W. C. Siegelranger, 42 *Waste in 15 min* 41G 31 (1947); *Chimie et Industrie* 35 (Heavy Industry Com) consisted of an elliptical-shaped 600-40-Arm nitrate TNT (bursting) rod with a cup section fitted with large fins (36 long charge), a long bonnet and a nose cone (10 long charge). Small, semi-day fins attached to the rear of the cup section fitted the surface was attached at the base. It is presumed that the bonnet provided with a large rod which fitted over the cup and
- 
- FUZE
- GAIN

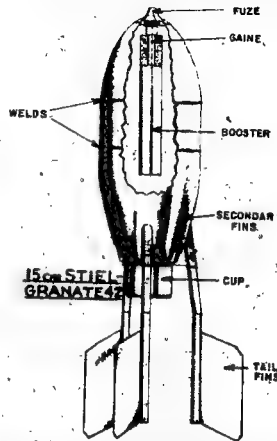
Rocket Signal Simulating Device (15 cm Maketen Schreib-
schuss Gerät, abbreviated as RSSG). See under Pyrotechnics
Antipathfinder Devices.

Rodded Bomb or Stick Grenade (Stielgranate). The following roddeed projectiles are described in TM 9-1985-3 (1953), pp. 108-110, 140-141:

- a) 3.7 cm "Spielmanns" 41 used in the Antitank Gun, 3.7 cm Pak 41 (Panzerabwehrkanone 41) consisted of an egg-shaped head (body) and a cylindrical tail provided with 6 fins. The head contained a shaped bursting charge consisting of 5.28 lb of 60/40-RDX/TNT (2 blocks wrapped in wax paper), two boosters (Kazdet).



a nose fuze (AZ 9075) and a base fuze (B4Z 5130). The tail portion of the projectile consisted of a rod which fitted into the bore of the gun, and a concentric perforated sleeve which fitted over the barrel of the gun. Tubular double-base NC-NG propellant (NGIRP), 6.61 oz. enclosed in a cartridge case, closed by two cost discs, was used as the propellant. Total weight of projectile 18.26 lb and overall length 27.362".



entered the muzzle of the gun before firing. This not dropped from the projectile about 150 yd from the muzzle. The bomb was propelled by 12.1 lb of propellant contained in a semi-fixed cartridge case. Total weight of the projectile was 105.0 lb and overall length 50.5". It was used against personnel had to clear minefields and wire obstacles.

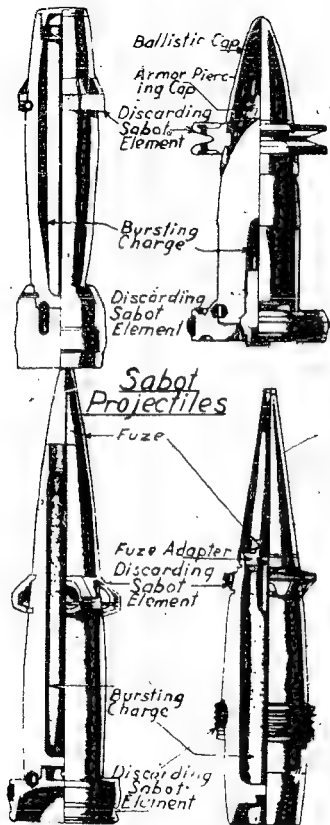
c) 37 mm Hollow Charge Stick Rifle Grenade, briefly described under Rifle Grenades, was similar to the 3.7 cm Stielgranate 41 (see also Stick Hand Grenade).

(See also Stick Hand Grenade)

Rebelsolninder Pulver (RZP). Finely pulverized iron prepared by atomizing molten cast iron by a cone of moist steam at a pressure of 2 to 3 atm. During this process most of the carbon is removed as CO_2 and thus removed. A large part of the iron was also oxidized during atomizing but it was recovered as pure iron on subsequent cooling in water and reduction with hydrogen. This powder was used in the manufacture of sintered iron items many of them of military use.

Reference: CIOG Final Rpt 595 (1945), p 52.

Röhrenpulver (RP) (Tubular Propellant). A propellant similar in form to the British Cordite. The compositions of some tubular propellants are given in Refs 1 and 2.



Salt (Säure) One of the older permissible explosives NG 11.2, colloidal cotton 0.5, Al nitrate 55.6, DNT 8.5, Na chloride 23.1 and carboxylic acids 2.5%; "Trenn" not value 287 cc and "Charge limit" 600 g.

Reference: A. Marshall, Explosives, London v 1 (1917), p 597.

Superoxide. See Nitric Acid.

Sänger-Bredt Missile, called Antipod Bomb, was a superoxide rocket designed by Dr. E. Sänger before 1942 but the project was abandoned without any practical development. This design embodied many unique features, which are briefly described by Göttsch on pp 57-8. It was planned to use the rocket in regions above a dense atmosphere. Each time it dived and hit a denser layer of air, the missile was supposed to bounce upwards. These movements would produce a kind of wave-shaped trajectory, similar to that obtained when a flat stone is ricocheted across water, but much less pronounced. As each plunge into a denser stratum would result in a partial loss of kinetic energy of the missile, the initially long jumps would gradually become shorter, finally to be transformed into an even gliding flight. It was presumed that this method would achieve a stable flight and a more accurate trajectory in a region above dense air, where conventional missiles usually behave rather erratically. The rocket was designed to be catapult launched and to be propelled by an oil liquid oxygen mixture, the calculated characteristics were: launching weight 220,400 lb, overall length (less booster) 91.8 ft, width of rectangular section 5.9 ft x 11.8 ft, maximum range 11,000 miles and maximum altitude 93 miles.

Reference: K. G. Göttsch, Development of the Guided Missile, "Flight" Publications, London, (1952) pp. 57-8 & 124-5.

Savin. See under Tillous.

Schwaben. An igniter using a compressed mixture of meal powder (Mealpulver) with a slow-burning substance such as a mixture of sulfur and K nitrate.

Reference: Kautsch, Chemische Untersuchung, (1944), p 555.

Sauerstoffkugel oder Sauerstoffpulver (Oxygen Balance or Oxygen Value). Abbreviated to O.B. it may be determined in the manner described in the general section or by the method given in A. Seestricher, Spreng- und Schießstoffe, Zürich (1948), pp 16-18.

Schulminomesschine. See under Kartmaschine.

Semina Pulver. One of the pre-WW II sporting smokeless propellants: gunpowder 95.0, TNT 4.0 and gelatinizer with "time moisture" 1.00.

Reference: H. B. Nussing, Das Schießpulver (1926), p 134.

Scheffler - Glück Fused Core, patented before WW II in America, was later improved and used as the Timedelay Fuse, D.K.G. It is briefly described in BROS Film Rept 644 (1945), pp 9-11. In Germany, this could replace the previously used Kronschmidt fused core "salvo-type" fuse.

Scheidtmann (Duke of Pickard Over). A mixture consisting chiefly of Ca and Mg nitrate was used during WW II in some substitute explosives (Ersatzsprengstoffe) as an extender of nitrocompounds which were not available during the war in sufficient quantity.

Reference: PB Rept 1420 (1945), p 11.

Schiesbaumwolle. See Schießwolle.

Schiesbecher. A rifled, caliber 30 mm, discharger cup which could be fitted to most types of German rifles. Was used for launching some anti-air rifle grenades. A photo of the Schiesbecher but no description is given in the Ordnance Sergeant, October 1945, p 9.

Rifle Discharger
Schiesbecher



This photo is by courtesy of Aberdeen Proving Ground, Maryland.

Schlesenschieber (Shooting Mortar). A device used for testing mines/explosives in galleries filled with fireclay and/or coal dust.

Reference: M. Lupus, S. Z. 26, 190 (1925).

Schleswille (Gucconin). Nitrocellulose of 13.2-13.3% nitrogen content, corresponding approximately to the Amer Gucconin. It was used in the manufacture of some smokeless propellants. (See also Nitrocellulose and under Propellants).

Schleswille (Schw) Explosives. See under Unterwasser sprengstoffe.

Schleswille 18 oder TSMV-101. An explosive described as Hexamino (Hexatin) in the general section. It consisted of TNT 60, hexanitrophenylamine 24 and Al powder 16% and was used in sea mines, torpedoes, depth bombs and underwater demolition charges.

Reference:

1) A. Seestricher, Procar 9, 33-41 (1943)

2) H. Munster, Procar 9, 62-65 (1945)

3) Allied and Enemy Explosives, Aberdeen Proving Ground, Md. (1946)

4) A. Seestricher, Spreng- und Schießstoffe, Zürich (1948), p 78.

Schloßpiste. (Sinking Distance). Same as Detonations-Bereichsgang.

Schloßpistenschere Sprengstoffe, oder Wassertsprengstoffe. Explosives safe for use in coal mines with fire damp. (See Wassertsprengstoffe, p 226 and also Schloßpistenschere Sprengstoffe)

Reference:

1) A. Seestricher, Schieß- und Sprengstoffe, Leipzig (1933), p 246

2) C. Heyling, A. Dreikopf, Sprengstoffe und Zündmittel, Berlin (1936), p 109

3) A. Seestricher, Spreng- und Schießstoffe, Zürich (1948), p 91

Schlaggesteinverwachsene, oder Verwachsene, (Hardpan Testing) Quarry. New options of galleries for testing explosives in regard to their suitability for use in various coal mines is given in the general section. The first German gallery was constructed in 1885 by Lohmann in Neunkirchen (Saarland). Other German galleries were: Drenze, near Herten, Gelsenkirchen-Saale, Gelsenkirchen and several galleries belonging to the plants manufacturing nitric explosives, such as Schleibach, Herten, Caspary etc. One of the newest galleries was in the Sächsischen Braunkohlrevier bei Freiberg (Sachsen).

Reference:

1) A. Marshall, Explosives, London, v 2 (1917), p 584

2) A. Seestricher, S. Z. 24, 788 (1929)

3) A. Seestricher, Schieß- und Sprengstoffe, Leipzig (1933), p 148-246.

Schmidlin G. 33 (SG 33). A rocket booster was invented by Schmidlin to increase the thrust of its 117 missiles thus assisting its take-off. (TM 9-1985-2 (1955), p 201).

Schnecken Pressen (Marm Press). In order to reduce the time of the rolling operation and to reduce the power consumption in the manufacture of solventless propellants, the Löbberg Fabrik of Dynamit A.G. rolls the NC-NG (or NL-DE-GUN) paste into pellets. The water content of this paste had previously been reduced to 8%, instead of 15-16%. It was used in the other propellant plants. In order to achieve such good dewatering, the usual centrifuging of the paste was followed by passing it through the Schnecken Pressen. The press consisted of a slotted barrel and an endless screw. When the paste was pressed some water escaped through the slots while the partially dehydrated paste was squeezed out ready for rolling into sheets (carpets).

Reference: A.A. Swanson & D.D. Sager, CIO Rept 29-24 (1946), p 7

Schnellmine. See Panzerschnellmine oder Landmine.

Schnellzünder (Quick Time Igniter), called also instantaneous Fuse and Quickmatch. Some German igniters, such as Demoschnitzler and Bichselzünder are described in Beyling-Dreikopf, Sprengstoffe und Zündmittel, Berlin (1936), p 229.

Schnitzel oder Schnitzel (Misapplied Non-German word) Schnitzel oder Schnitz, meaning Nose). The Dutch had fitted their submarines with an air intake back in 1940, and the Germans modified the device and called it Schnitzel. It consisted of a tube (about a dozen meters long), one end of which was connected to submarine Diesel, while the other end protruded above the surface of the water. The tube was divided lengthwise into two compartments - one for suction of air from the outside and the other for removing the gases of combustion of the Diesel. This device permitted the submarine to operate its Diesels while remaining in the submerged condition. In case of danger, the Schnitzel folded horizontally and the submarine submerged to a depth of at most 300 m (or even 400 m as was reported for the submarine Z1). As the material of the Schnitzel was usually non-metallic, it could not be detected by radar.

Due to the fact that the Schnitzel used during WW II did not supply an amount of air sufficient to replace the fuel air in submarine, it was necessary to replace the submarine after several hundred kilometers of underwater travel or equivalent duration. The maximum achieved in an uninterrupted submerged condition was 500 km.

References:

1. A. Ducrocq, Les Armes Secrètes Allemandes, Paris (1947), pp 20-24.
2. H. Schaefer, U-Boat 997, Noron, N.Y. (1950), pp 182-3.

Schoepher-Ringler Test. According to Shildon (Ref 1) this test was used in Germany to determine the suitability of cress paper intended for the manufacture of nitrocellulose. The Schoepher-Ringler Test was originally introduced into the paper industry to determine the freeness (viscosity) of the wood pulp. The test operates on the same principles as the Canadian Standard Freeness Tester (Ref 2).

1. L. Shildon, P.B. Rept 12,662 (1945).
2. J.M. Starcheson, Edit, Preparation and Treatment of Wood Pulp, McGraw-Hill, N.Y. vol 1 (1950), pp 943, 951 & 953 (See also Freeness and Test Testing in the general section).

Schoepher-Propellant. See Schoepher Shell.

Schoepher-Propellant (Schoepher). See under Landmines.

Schulze (Schulze) Type Igniter. Also called **Mohel-Schulze (Lover Type Igniter)** is briefly described under Igniters and in TM 9-1985-2 (1953), p 296. It was used in the Glanville 43 as an alternative to the Back Igniter.

Schulze Powder (Schulze Powder). An explosive patented in 1891. K. Schuler 60, pulverized antimony 25 and sugar 15%. A similar explosive was used by the British under the name **Schulze Powder**.
Reference: Daniel, Dictionnaire, Paris (1902), p 705.

Schulze Powder (Schulze Propellant). A smokeless propellant prop. about 1865 by Major Schulze of the Prussian Artillery, by allowing purified (degreased) wood (in the form of small equivalent pieces), followed by washing and boiling the resulting Nitrocellulose with water and then drying. After this the grains were impregnated with a concentrated solution of saltpetre, with or without Ba nitrate, and dried again.

Although this propellant was appreciably slower burning than earlier smokeless propellants consisting of outright compressed nitrocellulose (such as Von Schuler Propellant), it was still too quick for use in rifles, although quite suitable for shotguns.

Schulze propellant was manufactured not only in Germany but also in England (1868) and Austria (1870), but it did not achieve any success until it was modified in England by Griffiths and in Austria by Volkman. The Austrian propellant was made by partly gelatinizing the Schulze propellant with a mixture of ethereal oil and it became known as Celluloid. The British modifications began in 1885, compressed nitrocellulose wood pulp instead of previously used nitrocellulose. The composition of the British version Schulze propellant is given in Marshall (Ref 1), p 327.

The composition of German Schulze propellant given by Buewicz (Ref 2) was as follows: cotton 40, Ba nitrate 40, gun cotton 40, Ba nitrate 10, vaseline 8, moisture 1.3 and gelatinizer 0.5%.

1. H. Marshall, Explosives, London v 1 (1917), pp 47 & 127.
2. H. Buewicz, Das rucklose Pulver, Berlin (1925), p 134.

Schulze One of the Land Mines. See under Landmines.

Reference: TM 9-1985-2 (1953), p 178.

Schulze G. P. 40. Hollow charge m.f. granular described in TM 9-1985-2 (1953), pp 537-8. (See also under Rifle Grenades).

Schulze-Mineral. Same as Schulze-Min.

Schulze-Mineral (Black Powder). Composition, preparation and properties of black powders are given in the general section.

Table 58 lists some German military and commercial black powders.

Table 58
Black Powder

Designation	Composition, %		
	K nitrate	Charcoal	Sulfur
Geschützpulver, PPC/7	74.0	16.0	10.0
Colonne propellant 1871	76.0	15.0	9.0
Militärpulver 71	75.0	15.0	10.0
(Military rifle propellant 1871)			
Militärpulver (curtain)	75.0	15.0	10.0
Marine Geschütz Pulver	75.0	16.0*	9.0
(Navy Gunpowder)			
Jagdpatron (Hunting or sporting powder)	78.5	11.5	10.0
Sprengpulver (Blasting powder)	65.0	20.0	15.0
made by the Pulverfabrik	70.0	16.0	14.0
Spanisch	74.0	16.0	10.0
	66.0	21.5	12.5
Blasting powder	65.0	18.0	17.0
	(No al-)		
	76.0	14.0	10.0
Blasting powder B			

* Beech charcoal

References:

1. Gody, Traité des Matières Explosives, Namur (1907), p 71
2. R. Escalier, Schusspulver, Leipzig (1914), pp 160, 169 & 353
3. A. Schuler, Schulze's and Sprengpulver, Leipzig (1933), pp 97-112
4. E. Schuler, Química de los Explosivos, Madrid (1942), pp 275-9
5. A. Ammerbach, Spreng- und Schussstoffe, Zürich (1948), pp 59-6.

Schulze-Mineral. See Sulfuric Acid.

Schulze-Mineral Dynamite (Difficult Freezing Dynamite), called also **Unfreezing Dynamite (Non-Freezing Dynamite).** See Low-Freezing Dynamites in the general section.

Screaming Mini or Screaming Needle. According to H.H. Bullock of Picabury Arsenal, Screaming Mini was the nickname for any ammunition giving off a loud shrill sound in flight. One such item was the WW I 75 mm shell fired from the light, muzzle-loaded rifle mortar, called **Minerwerfer**. The shell had in the aft several vent holes that allowed air to pass through thus giving a shrill noise. Another well-known Screaming Mini was the 150 mm Smoke Rocket Projectile, 15 cm Rohbombe A/1, or its ammunition, used successfully during WW II. The

weapon, also nicknamed **Wack-Wack**, is briefly described in this section under **Rocket Launcher**.

(See also the general section).

1. W.B. Larson, Infantry Journal, September 1944, p. 23
2. Amos, Intelligence Bulletin, March 1945, pp 2-4.

See Dog. See Sechdog.

See Mother Bomb. See under Mother.

Securite. See Securite.

Securiphore. See Securiphore.

Sechdog (See Dog) (Chien de mer, in French). The "pocket" submarine (16 tons) with a radius of action of 500 km invented near the end of WW II. Its crew consisted of 1 or 2 men and it carried 2 torpedoes. It was provided with a small Diesel, generator, storage batteries, electric motor, oxygen tanks, and an arrangement which allowed it to submerge to as much as 30 or 60 m. This was an effective weapon which could do considerable damage if used in large numbers.

In addition to the Sechdog there were two other models of pocket submarines both propelled by electricity. The one, slightly larger than the Sechdog, was called **Seich (Schneider)**, while the other, considerably smaller, was called **Siber (beaver)**. (See also U-Boat, U-Boat).

Reference:
A. Ducrocq, Les Armes Secrètes Allemandes, Paris (1947), pp 31-3.

Seliger Sprengstoff. A permissible explosive patented in 1892 by Seliger of Berlin. It was made by blending 77 parts of K nitrate with 23 of the Na salt of sulphate/beta-nitrosulfonate, $C_{10}H_{10}SO_6Na$ [Daniel, Dictionnaire (1902), p 712].

Seliger Sprengstoff (Secondary Charge), called also in English **Base Charge**, **Heir Charge**, or **Lower Charge**, a charge in detonators or blasting caps which is placed underneath a primary or an intermediate charge. A secondary charge usually consists of a high explosive more sensitive to initiation than either P or TNT. The usual base charges were compressed nitryl, PETN, or RDX, while charges occasionally used included compressed P.A. and benzocinnonitril.

Securite (Securite). A type of mining explosive based on ammonium dichromate mixed with an oxidizer such as Am or K nitrate, patented about 1886 by F. Schuler.

Table 59 lists some securites.

Table 59

Component	Securite				
	1	2	3	4	5
Am nitrate	-	-	37.0	-	16.9
K nitrate	74.5	77.3	34.0	81.8	70.5
MNS with m-DNB	-	-	29.0	-	-
m-DNB	23.5	19.4	-	-	16.9
Am oxalate	-	2.0	-	3.0	-
Nitrocellulose	-	-	-	-	10.6

References:

1. J. Daniel, Dictionnaire des Matières Explosives, Paris (1902), pp 710-12
2. L. Gody, Traité des Matières Explosives, Namur (1902), pp 97 & 708
3. E. Colver, High Explosives, London (1918), p 141
4. F.M. Turner, Edit, Condensed Chemical Dictionary, Reinhold, N.Y. (1942), p 291.

Securiphore (Securiphore). A type of mining explosive moulded in Germany prior to WW I.

Table 60 gives some examples

Table 60

Components	Securiphore		
	1	2	3
Am nitrate	27.0	24.6	-
Ba nitrate	-	-	1.0
K nitrate	4.0	8.6	34.0
NG	40.0	36.4	15.0
Colloid cotton	1.0	0.9	-
Sebacic acid or its salts	12.5	11.6	-
No chloride	-	-	9.0
Rye flour	10.0	9.1	58.5
Wood meal	2.0	1.8	1.0
Liquid hydrocarbon	5.5	3.2	-
No carbonate or bicarbonate	-	-	0.5

References:

1. L. Gody, Traité des Matières Explosives, Namur (1902), pp 719-714
2. H. Marshall, Explosives, London, v 1 (1917), p 376.

Selbstzündende Probe (Spontaneous Ignition Test) for pyrotechnic compositions and their ingredients is described in **Kartillerie, Chemische Untersuchungen** (1944) 355.

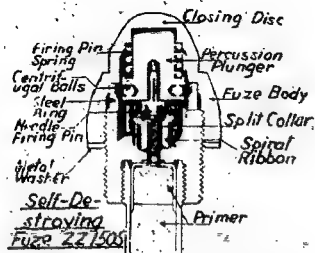
Self-Carrying Detonation Charge is described under **Kinemet** Factory, Dynamit A.G.

Self-Destroying Bullet. See Self-Destroying Train Bullet.

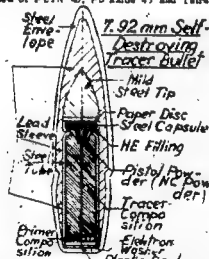
Self-Destroying Fuse, 22 1905, developed by the Deutsche Waffen- und Munitionsfabrik A.G., Liebert, was used in the 20 mm Mauser ammunition in air to ground flights. Like fuse AZ 1502 it was of the sensitive type required to function on a 2 mm paper screen at 100 meters. When the projectile was fired, the centrifugal force caused the steel balls to fly out into the enlarged portion of the retaining ring thus locking the percussion plunger and its compressed spring in place. The same force caused the brass spiral ribbon to unwind and increase in diameter until the shoulder; on the striker could pass through its center. By this time the projectile was a few meters away from the muzzle of the gun and the projectile was armed. On hitting the target the steel balls went back into their housings and the firing pin, activated by the compressed spring, pierced the primer cap.

If no impact took place within a range of about 2000 meters, the speed of rotation dropped to such an extent that the thrust of the balls against the metal surface was insufficient to support the firing pin spring. The primer was then fired and the projectile destroyed.

1. H. Perle, CIGS Rept 33-20 (1945), pp 69-70
2. Amos, TM 9-1985-2 (1953), pp 548-9.



Self-Destructing Tracer Bullet (Spitzgeschoss mit Zielkern, Leuchtgeschoss mit Zielkern) caliber 7.92 mm, developed during WW II by the Deutsche Waffen- und Munitionsfabrik A-G, at Lubek, was intended to be used against air or surface targets. It consisted of a steel casing containing a lead sleeve which enclosed a mild steel tip, a nose, capable with HE explosive filling and pistol powder, and a steel tube with tracer and primer compositions. The HE filling consisted of PETN 40, PB 40, and 20 and Tracer



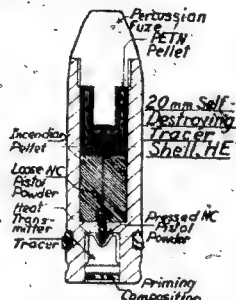
15%, whereas the pistol powder contained autocellulose with an ignition temperature of 160°. The bullet was self-destructed (in 500-600 m range), because the heat produced by the burning of the last portion of tracer composition ignited a small charge of pistol powder, which in turn set off the HE charge. The primer composition was ignited by the propellant in the cartridge.

Reference: H. Peplow et al., CIOB Rep 33-20 (1945), pp 28-9.

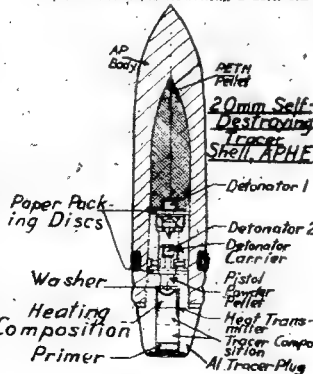
Self-Destructing Tracer Shells, caliber 20 mm, developed by the Deutsche Waffen- und Munitionsfabrik A-G, at Lubek, included the following:

a) HE Shell for Aircraft Guns. It was of conventional design and contained: a percussion fuse, a HE filling (PETN pellet), an incendiary pellet (Mg/Al alloy 50), a lead sleeve 40 and phenol formaldehyde resin 15%, a loose pistol powder (autocellulose), a pressed pistol powder, a heat transmitter, a tracer composition (two

increments, each pressed at 3500-4000 kg/cm²) and a pressing composition (pressed at 3200-3500 kg/cm²). If the shell was not exploded by the percussion fuse, it was self-destructed after about 0.3 seconds of flight. At this moment the flame from the last portion of the tracer ignited the pistol powder which in turn ignited the incendiary pellet. The intense heat produced by the burning pellet caused the HE charge to delaginate. The diameter of the tracer was 9 mm.



b) APHE Shell was of conventional design and contained a HE filling (PETN pellet), two detonators, a pistol powder pellet, a heat transmitter, a burning composition (the mixture 10.0, formaldehyde 36.0, in peroxide 22.5 and phenol formaldehyde resin 0.5%), a tracer com-



position (two increments) and a primer composition with its surface covered with NC lacquer. The shell was designed to give a trace of 4.2-4.8 sec duration, to penetrate a 20 mm armor plate and to explode 10-50 cm behind it. If the shell was not exploded in the above manner it was self-destructed by delagitation of the pellet caused by the intense heat produced on delagitation of the pistol powder, which, in turn, was ignited by the burning composition. This composition was incorporated in the shell because the heat produced by the tracer alone was not sufficient to ignite the pistol powder owing to the small diameter (6 mm) of the tracer compared with the diameter of the HE shell (9 mm).

Reference: H. Peplow et al., CIOB Rep 33-20 (1945), pp 34-61.

Self-Igniting Cushman. See Grankin, 1938.

Self-Propelled (SP) Gun Mount (Selbstfahrende (Sf or SB)) See under Panzer.

Servotop Gun. A mortar gun, caliber 800 mm, used electrically by the Germans during WW II at the siege of Sevastopol, Russia. The gun fired an 8 ton projectile with muzzle velocity of 2200-2400 ft/sec and maximum range of 29 miles. Weight of explosive was 2000 lb, wt of propellant 2500 lb, wt of gun 1375 tons, and length of barrel 169 ft. It is probable that the propellant charge was contained in a cylindrical casing made of a propellant composition, as described under Made-Up Charge.

Note: This gun was nicknamed Dorn or Gussak Gussak (See also under Weapons).

1) PB Rep 92 (1945), p 18.

2) Aberdeen Proving Ground, Museum; private communication.

Note: The projectile can be seen at the Museum.

SP Geschoss. See Spitzgeschoss.

Shaped Charge or Hollow Charge. See Hohlladung in this section and Shaped Charge in the general section.

Shredded Charge. See Maschenladung.

Shell. See Gussak.

Shell Mold Process or "Guss" Process of Precision Casting of Metals (Called also "Guss" Process or "Guss" Casting) developed in Germany during WW II by J. Croning, made possible the production of foundry molds and cores for cast metals in quantities and sizes that were formerly considered impracticable. In this process the thin shell-molds were formed by the adherence of a mixture, consisting of sand and plastics to heated metal pattern. Each shell mold was then hardened by further polymerization of the plastic of the pattern by heating for a short time in an oven with a pattern still attached. After removal from the oven, the molds were stripped from the pattern, clamped together in pairs in a box, lined with loose metal shot or other porous material, and filled with molten metal for casting.

The process is applicable to the manufacture of shells, bombs, grenades and rockets.

References:

1) J. Croning, Ger Pat Application No 48679 (1949), described in PB Rep 8393 and 81284.

2) H. H. Ames et al., The Formator, August 1950, pp 92-95 and 206-17.

3) H. L. Day, The Iron Age, 149, 28 (Jan 1952).

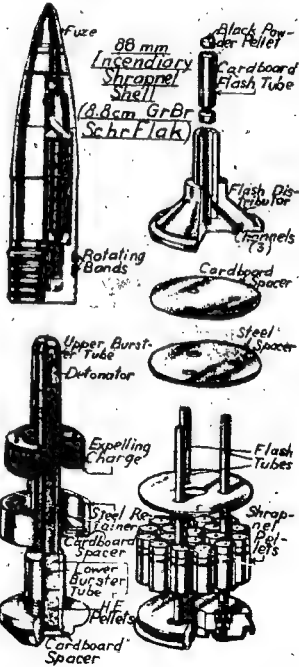
4) R. H. Ames et al., The Formator, June 1951, pp 112-17 and 281-95.

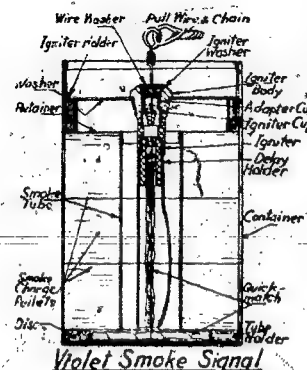
5) R. H. Ames, PB Rep 106640 (1952) (47 references).

Shrapnel or Shrapnel Projectile. See Jagdgesch.

Shrapnel Mine (Schrapnellmine, abbreviated as S-Mine, sometimes called Schitzmine). Two types, S-33 and S-44, are described in TM 9-108-2 (1951), pp 279-81. Owing to the fact that three mines rose into the air (to the height of 1 to 5 feet) before exploding, they were nicknamed Bouncing Mines (See under Landmine).

Shrapnel Projectile (Schrapnellgranate). Only one such projectile, namely the 8.8 cm German Brand Schrapnell (8.8 cm Brand Schrapnell) is described in TM 9-108-2 (1951), p 440-49. The projectile consisted of a thin steel shell of conventional design containing: 72 incendiary pellets, a point detonating





Violet Smoke Signal

assembly, which included the delay element, was extended from the igniter holder. Below the igniter, inside the central tube, were located loose pieces of quickmatch (black powder) used to facilitate the ignition of the delay element.

The signal was fired by removing the cover, pulling down on the igniter chain (by means of the pull ring) and then throwing the signal (or placing it upright on the ground). The friction was being pulled through 0.04 g of the composition; and more sulfide 70, potassium chlorate 30 and mercury (fulminate 20%, covered it to flash and to ignite in turn, the delay element (0.05 g of K nitrate 75, charcoal 15 and sulfur 95%). After burning for about 1/2 seconds, the flame from the delay element ignited one cord of quickmatch (black powder) which, in turn, ignited the black powder composition (1.3 to 1.8 g) in the bottom igniter disc and finally the smaller amounts of the smoke from the burning charge was forced through the small hole in the central tube (or in the central cavity in the case of the green signal), and thence around the friction igniter, and through the hole in the remnant into the space between the remnant and igniter holder. The pressure of gases generated on burning ruptured the film covering the six vent holes in the igniter holder thus allowing the smoke to escape. It was assumed that the smoke charge burned from the center outward and from the bottom upward. The duration of emission of smoke was 12 to 20 seconds.

Signal Smoke Device. See Signal Device.

Silene Sprengstoffe (Silene Explosives) were chlorine explosives developed under WW I by the Oberheftungs A-G für Fabrikation von Lignos (Schleusen- und Schiffbau für Armee und Marine). According to Einsaite (Ref. 1), p. 193 one type of Silene was a mixture of K chlorate 80 (max) with 20% resin of which 4% could be in the nitrate state. Another composition contained K chlorate 75 (max) resin (amine) and Na chloride 10% (min). The resin had a m.p. of about 70, and the Na chloride was mixed with 10% of its weight of paraffin oil.

Following were the compositions of some of these explosives:

- Silene IV: K chlorate 70, resin 6 and Na chloride 22%. It was suitable for blasting rocks and ore, but could not be used in gaseous or dusty coal mines (Ref. 1).
- Silene No 4: K chlorate 80 and resin 20%. It was suitable for blasting rocks and ore, but could not be used in gaseous or dusty coal mines (Ref. 2).

References:

1. R. Einsaite, Chlorosprengstoffe, Velt, Leipzig (1910), pp. 143 & 185.
2. Admittable Explosives, Churchill, London, v. 1 (1917), pp. 382-3.
3. E. Baran, Explosives, Van Nostrand, N.Y. (1919), p. 111.

Silver Azide (Silberazid) (Ag Az). See general section under Azides.

Silver Fulminate (Silberfulminat). See general section under Fulminates. It was used in Germany as a primary charge in the Amant-Hausen (q.v.).

Silver-oxide Fiber (Silberoxy-Faser). A type of blasting explosive, made by mixing powdered pyrite dust (left over from WW I) with 5 to 10% of aqueous solution of cellulose pitch, a waxy product obtained by evaporating silicic acid from the pulp industry. The composition could contain up to 10% of aromatic nitrocompounds such as TNT, DNAB, etc.

References:

1. P. Nardin, Schaefer and Sprengstoffe (1927) p. 66.
2. J. Perin, Lethal, Powder, etc. (1935), pp. 457-8.

Silverydust oder Sponyd. Primary explosive mixture developed in Germany about 1930 to replace previously used mercuric fulminate compositions. It has been claimed that the production of decomposition of Silverydust was non-explosive and did not involve first-order reaction. Kovachek (Ref. 3) gave the composition of a mixture used by the Germans during WW II as follows: lead azophane 25 to 55, urethane 1.2 to 1.6, nitrobenzene 25 to 45, PbO₂ 5 to 10, Sb₂O₃ 5 to 10, Ca sulfide 3 to 25 and powdered glass 0 to 5%.

References:

1. E. von Herz, S.S. 28, 39 (1933), Die erfindungsfähige Zündung.
2. A. Schuchter, Spreng- und Zündstoffe, Zeitsch. (1948), pp. 98 & 106-7.
3. H. Fickert, A. Kovachek, Min. post. 31, 26-27 (1949).

Showered Iron and Steel Items, such as bullets, pyrotechnic devices, etc. are mentioned under Pyrotechnics.

Showered Iron Protection. See under Tinfluoride Verfahren.

Ship Bomb or Kort Apparatus, designated as SB 400 Kwai K in description on p. 14 of TM 9-1935 (1935). (See also under Bombs).

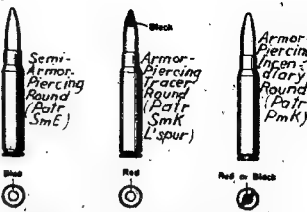
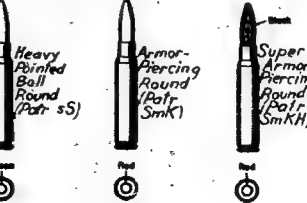
Small Arms (Handfeuerwaffen). See under Weapons.

Small Arms Ammunition. According to A. J. Dore, Ordnance Sergeant, December 1943, pp. 357, the German small arms ammunition was similar to American. The complete round consisted of a cartridge case, percussion cap (primer), propellant charge, and bullet. The cartridge was drawn either from sheet brass (copper 71 and zinc 28%) or from sheet steel, copper plated on both sides. The case was bottle shaped, grooved at the base and coned slightly to facilitate extraction. A primer pocket was formed in the base of the case and was connected to the interior by flash channels. In the center of the pocket an anvil was formed on which the primer composition was fired by the firing pin. The primers were of the Berdan type, either the No 88 or No 10. The No 88 primer consisted of a brass cap containing the primer composition, and a covering cap of double-size zinc-plated lead foil. The primer composition was put into the cup dry and was protected from dampness by the cap which was lacquered on the inside. The inside of the cup was also lacquered to the level of primer composition. The No 10 primer was essentially the same as the No 88 except that its primer composition was different and practically non-erodable. A charge of a typical small arms cartridge consisted of a single-base (nitro-cellulose) propellant in blackish, coarse, granular-shaped flakes about 2.2 mm thick and 1.2 to 1.5 mm long, with smooth-cut surfaces. A typical bullet had a boat-tail base and consisted of a lead core and jacket consisting of either cupronickel, gilding metal or copper-plated steel. There were also bullets with steel cores or made entirely of steel (See under Steel and Iron Ammunition Items). The bullet was crimped to the cartridge case in the conventional manner by means of a cannelure.

The following calibers were commonly used during WW II:

A. 7.92 mm Ammunition which can be subdivided into the following types:

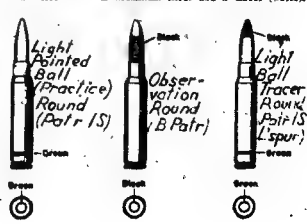
- Per all (Patronen schweizer Spitzgeschoss), Heavy Pointed Ball Ammunition, had a bullet with a lead core and a copper jacket. The bullet was painted green, the base of the cartridge was painted green, if labeled as simply Pat all, the ammunition could be used in rifle in either (as in Manner of Gewehr 41) or in machine guns (such as MG 15, MG 17, MG 81, MG 14 and MG 42). In the machine gun it was used as a round ammunition with label "Pat all" in which the letters "MG" indicated that this round was cap packed. The label "Pat all" for Gewehr 41 indicated that the round was designed for use in rifle and the label "Pat all" for MG indicated that the round was designed for use in machine gun.
- Per SmK (Patronen Spitzgeschoss mit Stahlkern), Armor-Piercing (Super) Ammunition, had a bullet with a longer than in (a). The core was of steel and the jacket of steel with gilding metal coating. The annulus was painted red.

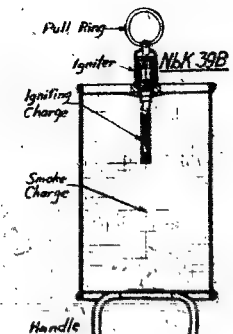


- Pat SmKH (Patronen Spitzgeschoss mit Stahlkern), Gilded, Armor-Piercing (Super) Ammunition, had a bullet with a tungsten carbide core and a steel jacket coated with gilding metal. The bullet was painted black and the annulus was red.
- Pat SmE (Patronen Spitzgeschoss mit Eisenkern), Semi-Armor-Piercing Ammunition, was similar to the above, except that the case was of soft steel or iron. (See also under Steel and Iron Ammunition Items).
- Pat SmK (Patronen Spitzgeschoss mit Stahlkern), Heavy Pointed Ball Ammunition, had a bullet with a lead core and a copper jacket. The bullet was painted black and the annulus was red. This round was used principally against aircraft.
- Pat SmK (Patronen Spitzgeschoss mit Stahlkern), Heavy Pointed Ball Ammunition, had a bullet with a lead core and a copper jacket. The bullet was painted black and the annulus was red. This round was used principally against aircraft.

Note: This bullet is described more fully under Observation Bullet. According to CIG Report 3-20 (1947), p. 3, it was also adapted as an incendiary bullet for use against ships.

- Pat SmL (Patronen Leichter Spitzgeschoss mit Leuchtspur), Light Ball-Tracer Ammunition, had a bullet with an aluminum filler and a tracer (which was painted red).





39.5 and handle 1.0%. After burning for about 3 seconds, the smoke charge was ignited. The smoke and gas pressure on burning forced an exit through the zinc top layer beneath the two holes in the steel top. A large volume of dense gray smoke was emitted, according to CIOB Rept 32-13 (1945), for about 1 minute or 4-7 min according to P-47H 140.

d) Fast Smoke Candle (N6K 39B) was similar in construction to the N6K 39B with the exception of the filling and the method of use. Its smoky mixture consisted of Hexa 47.5, Zn dust 47.5 and Ba nitrate 1.0%, compressed to form a 4" cylinder weighing 1.7 kg and having a burning time of 100-200 sec. It was operated by firing from a projector attached to a cable.

e) Slow Smoke Candle (N6K 142) consisted of a round, sheet metal container about 480 mm long and 100 mm diam, with a compressed increment of smoke composition (Hexa 65, Zn dust, coarse 27, Zn dust fine 10 and Ba nitrate, added 0.75-1.750 weighing 17.3 kg. It was ignited by means of a 300 g layer containing Hexa 47.5, Zn dust 47.5 and Ba nitrate 1.0%. The emission time was 15-25 minutes.

f) Black Smoke Candle (N6K 142A) was identical in construction with the previous candle but contained a different smoke composition: Hexa 28, K chlorate 38, crude antimony 10 and Hexamethyl 1%. It was operated in three variations, total weight 12.15 kg. Ignition was effected by means either of a safety fuse igniter or a low tension electric igniter and a glass. The time of emission was 10-16 minutes.

g) Smoke Candle (N6K 55.6) which served as a fixed "artificial" smoke marker, consisted of a sheet metal cylinder, 140 mm long and 91 mm diameter packed with 20 mm diameter emission holes and filled with a compressed mixture of Hexa 12.5, Zn dust 38.0, ZnO 40 and Mg powder 1.5%. Ignition was effected by a 60 mm long fuse and a glass. The time of emission was 45-75 seconds.

h) Black Smoke Candle (N6K 55) which served as a fixed "artificial" smoke marker, consisted of a sheet metal cylinder, 140 mm long and 91 mm diam, provided with 20 40 15 mm diameter emission holes and containing two compressed pellets (total weight 1.2 kg) of a mixture of Hexa 7, K chlorate 45 and crude antimony 50%. Some ignition assembly as above. Time of emission about 2 minutes.

a) Black Smoke Candle (N6K 55) constructed from a peashooter 3 mm thick, was of the same dimensions as the above sheet metal container. The filling consisted of two compressed increments (total weight 1.2 kg) of Hexa 7, crude antimony 50 and Mg powder 140%. Time of emission about 2 minutes.

b) Smoke Candle (S6K 11) which served to simulate the burning of vehicles, consisted of a gastight cylinder, 56 mm diameter and 190 mm high, filled with two hand pressed increments (total weight 400 g) of a mixture: Hexa 28, K chlorate 40 and crude antimony 32%. Time of emission of black smoke "vehicle" "burning" indicator, consisted of a sheet metal tube about 700 mm long and 80 mm diam. The smoke mixture consisting of Hexa 40, Zn dust 47.5 and Ba nitrate 1.0% weighing 12 kg, was pressed in directly. Ignition was effected by a low-tension fuse and a glass. The time of emission was about 10 min.

Smoke Compositions (Reactants). Smoke compositions may be subdivided into two types:

a) Compositions which on burning developed a dense white or black smoke serving for screening purposes (Nebelstift).

b) Compositions which on burning developed a colored smoke (Donnuch), serving for signaling purposes. Many of these compositions are denoted under signal devices, smoke bombs, smoke candles, smoke generators, smoke projectiles, smoke signals and under pyrotechnics.

According to CIOB Rept 32-13 (1945), p. 18, several smoke "compositions" were developed for use at the end of the VB II but were never put into service. Several compositions were prepared by adding to the mixture of Hexa (hexachloroethane) and Fe powder varying amounts of Mg, to accelerate the reaction. One such mixture contained Hexa 55, Fe 15 and Mg 7%. Very effective fumigants giving rise to orange smoke, were obtained by varying the proportions of the composition Hexa 48, Fe 10, Mg 36 and Mg powder 16. A new mixture designed for smoke candles contained Hexa 55, Zn dust 40 and ZnO 10%.

Among other smoke compositions may be mentioned titanium tetrachloride, designated as FM (used in some smoke hand grenades), a mixture of oleum 80 and pure zinc 20% (used in some projectiles) and a black smoke mixture (Mg 15.5, hexachloroethane 6.5%, methanol 12.0 and antimony 1.0% (used in the Black Smoke Candle)).

1) E.V. Bauman, CIOB Rept 32-13 (1945), pp. 10-18
2) L.H. Spill, CIOB Rept 32-16 (1945), pp. 17-18
3) L.H. Spill, CIOB Rept 32-17 (1945), pp. 12-14
4) Amm. TM 9-1978 (1953), pp. 402, 417, 497 & 506.
(See also references under Colored Smoke).

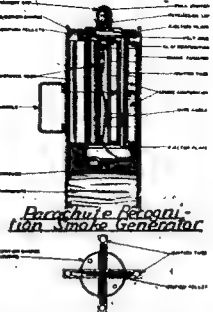
Smokeless Propellants or Smokeless Powder (Rachkono Pulver oder Rauchkornlose Pulver). See Propellant.

Smoke Flare. See under Flare.

Smoke Generator (Rauchentwickler). According to E.V. Bauman, CIOB Rept 32-13 (1945), p. 10, all German generators examined by him consisted of a sheet metal container with one or several "emission holes" filled with one of the varieties of Berger's mixture. In these mixtures the hexachloroethane was dissolved. All the containers were used as the source of chlorine and zinc reacted with metals such as Zn or Fe. The metal melt then carried away smoke of any desired color was desired. All the containers were ignited by means of an igniter assembly.

Several models are described in this German section under Smoke Candles. They are listed in German Nebelkorn.

One of the generators, namely, *Propellier-Entwickler* Smoke Generator is described in TM 9-1978-2 (1953), pp. 89-92. The device consisted of an aluminum cylinder divided into two parts, one housing the smoke composition and the other the peashooter. The first section was subdivided into sub-sections by means of a series of cones connected by revolve metal distance rods. Eight of these rods were equally spaced around the circumference of the plates while the remainder four were spaced at distance from and closer to the center. The smoke cylinders



were finally held in two turns, each with four cylinders. Four 1.4 in holes were drilled in the plates for the igniting tubes. The individual smoke cylinders were aluminum cylinders filled with acid waterproof paper and containing four smaller blocks, three of smoke composition and one of a clay-like substance. The smoke composition consisted of a bent stable blue dye 48 mixed with K chlorate 35, lacquer 7%. Each of the three smoke composition blocks had a small quantity of priming composition (black powder) placed in the loose connection at the base before passing to master ignition between one block and the next. The ignition pellets were arranged to accept the flash from the reaction charge and distribute it to the four igniting tubes, each of which pierced the center of two smoke cylinders. A total of fourteen black powder ignition pellets were packed in these tubes. The reaction charge, positioned directly below the pull igniter, consisted of 55 g of fine grained black powder. Below this was the first metal electric plate which had a hole in the center to allow the flash to reach the igniting pellets. The second electric plate, designed to prevent the reaction from burning, was arranged as a ring in the outer container, was placed in the lower part of the upper container directly above the plate. The peashooter canopy was made of continuous filament viscose rayon. Total weight of the generator was 27.5 lb, overall length 20" and maximum diameter 9".

For operating the device, the metal cap was removed, the friction igniter cap was unscrewed and pulled longitudinally, and the reaction allowed to fall clear. After a delay of 4 or 5 seconds, the igniter (located over the flash from the detonator) passed to the reactor charge to explode it. The pressure of the gases of explosion forced out the upper (small) section of the cylinder which, in turn, pulled out the peashooter. The peashooter was fired from the reaction charge against the pellets of black powder which distributed the flame to the four igniting tubes, each of which pierced the center of two smoke cylinders, thus igniting the smoke composition. Each cylinder emitted smoke of good density for about 26 seconds.

Smoke Grenade. See Smoke Hand Grenade and under Pistol Grenades and Rifle Grenades.

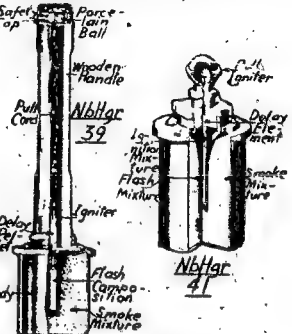
Smoke Hand Grenade (Nebelhandgranate oder Blendkörper). The following types are described in TM 9-1978-2 (1953), pp. 317-330:

a) Smoke Hand Grenade 39 (N6K 39), commonly known as the HE 39, was a 3.5 in diameter (external) and 4.7 in long and 1.7 in diameter filled with a smoke composition. It was filled with a smoke mixture containing hexachloroethane and Zn dust. Total weight 1 lb 14 oz

and overall length 14". Duration of smoke 2 minutes. It was used for screening machine gun nests and pull boxes (p. 3307).

b) Smoke Hand Grenade 41 (N6K 41) was similar in construction to the N6K 39, except that it was not provided with the stick (handle). Maximum diameter 4.7", overall length 4.7" and total weight 1 lb. was filled with hexachloroethane - Zn dust mixture. Same time of emission as in the N6K 39 (p. 32-63).

Note: According to CIOB Rept 32-13 (1945), p. 11, the composition of the smoke mixture was: Hexa (hexachloroethane) 55.0, Zn dust 43.5, and Ba nitrate 1.0%. The weight of the charge 400 g and the time of emission 150-250 seconds.

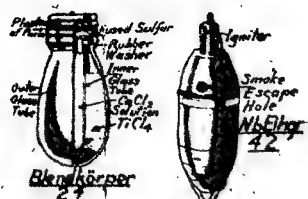


c) Smoke Hand Grenade (Blendkörper 14) consisted of a rear door glass plate (15" diameter), provided with a cardboard handle and filled with 100 g of titanium tetrachloride (FM). Its overall length was 6" and total weight 3.5 lb. The reaction was used to produce a small smoke screen to blind the enemy or to mask gaps in larger smoke screens. The flash could be extinguished by covering the reaction with a surface. On vaporization the tetrachloride formed a dense smoke, if the relative humidity was high (p. 327-8).

d) Smoke Hand Grenade (Blendkörper 24) consisted of a cover glass plate 14" (diameter) containing 270 g of titanium tetrachloride (FM) in a sheet glass plate containing 36 g of an aqueous solution of Ca chloride - which was sealed on a rubber washer in the back of the outer container. The ensemble was sealed by a valve and cement plug. The contents of the inner tube served to provide the water necessary for the reaction with tetrachloride in the formation of heavy smoke. The Ca chloride was quickly added to the water. The grenade was operated in the same manner and for the same purposes as the Blendkörper 14 (p. 328).

e) Egg Type Smoke Grenade (N6K 42) consisted of a cylindrical-ellipsoidal shaped metallic container, 4.9" long and 1.7" diameter filled with a smoke composition. One end of the body was flattened to permit the insertion of the pull rope igniter ZS-Schuss 9 (p. 329).

Note: According to CIOB Rept 32-13 (1945), p. 13, the composition of the smoke mixture in the N6K 42 was: Hexa (hexachloroethane) 55.0, Zn dust 43.5 and Ba nitrate



1. The weight of the mixture was 170 g and the time of initiation 60-100 seconds.

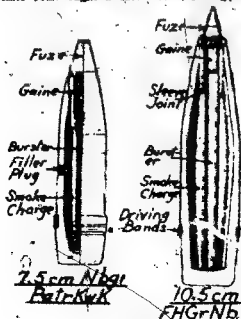
Smoke Signal Device. See under Signal Device and also under Pyrotechnics.

Small Signal Groups. See under Pistol Grenades.

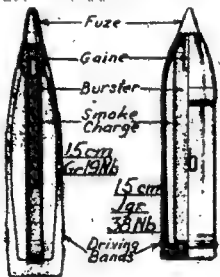
Smoke Projectile or Shell (Nebelschnecke, Rauchgranate). Projectiles containing a large charge of smoke producing composition and a small charge of bursting explosive. Several types of such shells were used during WW II by the Germans. These shells, on explosion, produced much smoke which was effective against personnel (not on objects) and a dense smoke or fog which served to prevent the enemy from seeing that was going on. In some cases the smoke projectiles were used for signaling purposes, as for instance the 80 mm Colored Smoke Signal Projectile.

The following smoke projectiles are described in TM 1-1965-1 (1955), pp 402-3, 472-3, 496-8, 506-7, 512 and 513-7:

a) 75 mm Smoke Projectile for the Tank Gun (7.5 cm Wgr) (see 6.02) was designed to the most design at the HE Projectile. The inner tube contained a small burning charge (2 cm of picric acid) and a large charge of smoke. 80 mm smoke projectiles were used for signaling purposes. Total weight of shell was 13.6 lb (pp 102-3).



b) 105 mm Smoke Projectile for the Fish Howitzer (10.5 cm FHGrNb) was similar in construction to the previous shell. It contained 4.3 lb of P.A. (burning charge) and 4.1 lb of smoke charge (oleum impregnated in pumice). Total weight of projectile 10.5 lb (pp 472-3).
 c) 150 mm Smoke Projectile, Type 19 (15 cm Wgr 1944) for the Heavy Howitzer (15 cm Wgr 1944) or Wgr 18, it contained 1.21 lb of P.A. (burning charge) and 14.0 lb of oleum impregnated in pumice. Total weight of projectile 35.8 lb (pp 496-8).
 d) 150 mm Smoke Projectile (15 cm Wgr 1944) for the Heavy Infantry Gun (15 cm Wgr 1944) it contained 1.21 lb of P.A. (burning charge) and 14.0 lb of oleum impregnated in pumice. Total weight of projectile 35.8 lb (pp 496-8).
 e) 150 mm Smoke Shell, Type 18 (15 cm Wgr 1944), for the heavy Field Howitzer (15 cm Wgr 1944) was similar in construction to the 15 cm Wgr 1944, except that its burning charge consisted of TNT. Total weight not given (pp 506-7).



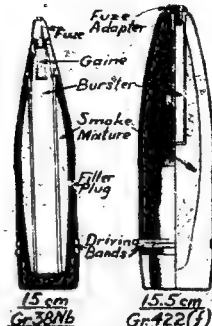
f) 155 mm Smoke Projectile (15.5 cm Gr 422 (P) for the French Heavy Gun 15.5 cm Gr 422 (P) Lisle 1916 St Ch was of conventional design. Its inner (center) tube was shorter than in the German designed smoke projectile and extended to less than half of the length of the shell (p 512).
 g) 80 mm Smoke Projectile (8 cm Wgr 1944) for the medium (medium) gun (8 cm Wgr 1944) and for the short (short) gun (8 cm Wgr 1944) was conventional in design. It carried a small burning charge of smoke mixture and a 2.75 cm Wgr burning charge. It weighed 7.85 lb and was provided with 12 lines (p 512).

h) 80 mm Cannon Smoke Projectile (8 cm Wgr 1944) (see 6.02) was of conventional design and carried 12 lines. It was filled with a composition which gave a colored smoke on burning (p 511).

i) 300 mm Smoke Projectile (30 cm Wgr 1944) (see 6.02) was of the same design as the corresponding HE mortar projectile described on p 515 of TM 1-1965-1 (1955).

j) 1915 mm Antitank Projectile (25.5 cm Gr 424 for the Howitzer (25.5 cm Wgr 1944) is briefly described under Spontaneous Projectile.

k) 105 mm Field Howitzer Smoke Shell (10.5 cm FHGrNb) (see 6.02) was described on p 14 of CIGOS Repts 32-13, (1945) was filled with 1.8 lb of the smoke mixture



containing: hexachloroacetone 55, Zn dust 43.5 and Be nitrate 1.5%. The time of emission was 4-7 minutes. Note: According to H.R. Patrick of Picatinny Arsenal, all German smoke and chemical projectiles were loaded from the side. This was contrary to the American practice of loading projectiles through the nose.

Smoke Pull Charge. According to H.J. Eppig, CIGOS Rept 32-56 (1945), this device was developed by the Deutsche Pyrotechnische Fabrik at Kieselbach/Vacha, but the item is not described.

Smoke Rifle Grenade. See under Rifle Grenade.

Smoke Rocket. See Smoke and Chemical Rocket.

Smoke Shell. See Smoke Projectile.

Smoke Signal Device. See under Signal Device.

Smoke Signal, Main. See under Signal Device and also under Pyrotechnics.

Smoke Stick (Nebelstange), which served as a wind direction indicator, consisted of a sheet metal tube, about 100 mm long and 16 mm diameter, attached to a wooden handle about 10 mm long. The smoke filler consisted of six pellets containing: lactone, K chlorate and Am chloride (react composition is unknown). It was ignited by means of a cup with a friction surface.

Reference: E.W. Batesman, CIGOS Rept 32-13 (1945), p 18.

Smoke Tube (Rauchrohr), was a smoke emitting device consisting of a seamless drawn tube, 250 mm long and 25 mm diameter, into which the following compositions were pressed by hand:

a) 1st layer: hexachloroacetone 49, Zn dust 41, Zn oxide 4 and Mg 05 and
 b) 2nd layer: hexachloroacetone 55, Zn dust 41 and Mg 45. Ignited by a wick from the rear of the tube. Total weight of the device was about 200 g and time of emission not less than 50 sec.

Reference: CIGOS Rept 32-13 (1945), pp 11-14.

Smoke Type Igniter (Rauchkinder). See under Igniter.

Smokeless Gun. See under Gun.

Smokeless Gun. An explosive containing Na nitrate 55 and TNT 45%. It was usable for loading bombs and shell shells. (A. Stenpecker, Schiene- und Sprengstoffe, Barth, Leipzig (1933), p 277.)

Sodium Azide (Na Az) (Natriumazid). See general section under Azides. Na Az was used in Germany for the manufacture of lead azide (L.A.) as described in WD Rept 93,613 (1947), Section D (See also under Blisidat).

Sodium Chloride Explosives or Kitchen-Salt Explosives (Kochsalzexplosive). German substitute explosives containing large amounts of NaCl (up to 60%). They are described under Explosives.

Sodium Nitrate Explosives (Natriumnitratexplosive). Explosives containing Na nitrate, such as Sodanox and some explosives described under Explosives.

Sodium Picrate (Natrium Pikrat). See general section under Picrates. It was used during WW II in Germany as a component of GP (Powder), prepared as a substance for black powder and as a propellant for Panzerfausts. In this composition the picrate was mixed with a binding substance such as latex SS.

Reference: CIGOS Rept 25-18 (1945), pp 27-28.

Solid Catalyst. See M-14.

Solvents and Plasticizers for nitrocellulose, plastics (such as polyvinyl chloride), resins, synthetic rubbers etc were described in some CIGOS and FIAT Reports, and especially in CIGOS Repts 1651 and 1652. These two reports covered the investigation during November-December 1945 in the field of solvents and plasticizers sponsored by the Raw Materials Division of the (British) Board of Trade. The field of investigation did not include petroleum and chlorinated hydrocarbons. A brief description of the methods of preparation of about 150 solvents and plasticizers were given, but no data for the properties of plastics are given in the above reports.

Somax. See under Tolilons.

Sondermunition (Special Propellant Material), developed during WW II by IG Farben, was presumably intended for use as jet propulsion fuel. It contained an unsaturated compound (diisobutylene) which reacted with concentrated (90%+) nitric acid with explosive violence. The reaction time was within hundredths of a second.

The mixture finally developed contained divinylacetylene (diisobutylene), 5-6, vinyl acetate 6-12, benzene 70, 1 and 10 (vol/vol/vol/vol).

Note: The composition does not add to 100%. The large amount of iron catalyst appears questionable. Reference: CIGOS Rept 25-18 (1945), pp 20-21.

Sound Gun. This weapon, constructed by R.Y. Altschuler of Austria, was designed to cause casualties or damage by means of sound waves of great intensity. It was claimed that at great range (say 60 m) it could kill a man and at appreciable length of time, A. brief description of this device is given by L.E. Simon, German Research in WW II, Wiley, N.Y. (1947), pp 181-2. The weapon consisted of a parabolic reflector, 3.5 meters in diameter, having an attachment extending to the rear of the vertex of the parabola. The attachment consisted of a firing chamber (for propellant) and a sound horn, the length of which was of the wave length of the sound. At its rear, the chamber was provided with two conical nozzles, the outer nozzle emitting methane and the inner one emitting oxygen. The frequency of sound was from 800 to 1500 impulses per second and the pressure produced by the sound waves was

Ger 187

Spezifische Energie oder Spezifischer Druck, designated as "E". See Specific Energy, or Specific Pressure in the general section.

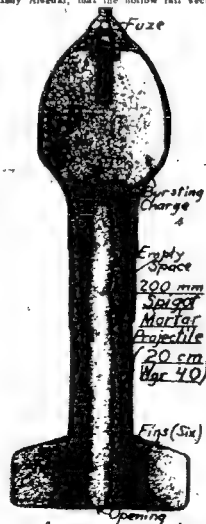
Spezifisches Gewicht See Specific Gravity in the general section.

Spezifische Wärme See Specific Heat in the general section.

Spigot Motor (Lanzengewehr) Projectile. The following projectiles are described in TM 9-1985-3 (1953) pp 334

- 200 mm Motor Projectile, 20 cm Wgr 40 (reference 40) for use in the light (lighted) spigot motor 20 cm (Lanzengewehr) consisted of two sections, one housing about 1 1/2 lb of bursting charge (TNT) and the other projectile in other sections each weighing 12 g. Total weight of the round was about 50 lb (p 334)
- 380 mm Motor Projectile (38 cm Wgr 40) for the heavy spigot motor (38 cm (Lanzengewehr) was similar in design and shape to the 20 mm projectile. It contained 110 lb of HE bursting charge and was provided with 6 fins. Total weight of projectile was about 328 lb (p 335).

Note: There is no indication in the above manual how this projectile was fired and what kind of spigot motor was used. It is probable, however, according to H.H. Bullock of Picatinny Arsenal, that the hollow tail section of the



equal to 1000 microbars, when measured at a distance of 50 meters. The military value of this weapon was slight due to its short range.

*Some Guidance System for Missiles. See under Guidance Systems for Missiles.

Spore Explosions with Carbon Dust. See under Kilmoril Fabrik Dynamit A-G. Preparing of Explosives and Research and Development Work.

Spall Faceheads or Splitting Priming Drops. When shooting in coal mines where considerable uncontrolled electric currents are to be found, the faceheads of electric blasting caps or detonators have to be constructed in such manner that they shall not ignite from a potential as high as 15 volts. This was achieved at the Troisdorf Fabrik, D.A.G. by using special heavier faceheads in the resistance range of 1000 to 10000 ohms.

For preparing such faceheads the tip of the bridge wire was dipped successively into the following compositions, allowing the material to dry after each dip:

- 1st dip composition, which consisted of Pb peroxide 43 g, carbon - magnesium alloy 28.5 g and Al (particle size 10 to 20 microns) 28 g suspended in about 70 ml of a 1% solution of NC in acryl or butyl acetate.
- 2nd dip composition consisted of red lead (particle size less than 5 microns) 90 g and silicon (particle size 20 to 40 microns) 10 g suspended in a 1% solution of NC in acryl or butyl acetate.
- 3rd dip composition was a lacquer consisting of a 15% solution of NC in 75.25% butyl acetate/dioxane, to which was added 5g of ADM (methylcyclohexyl ether of adipic acid) in the amount of 10% of the dry weight of NC.

The average stability of these faceheads in moist atmosphere was not very good.

Note: Soldering of the bridge (base) wire to the lead-in wires necessitates the use of magnesia for facehead dips, preparation of NC lacquers and the process of dipping the facehead combs very described under Facehead Manufacture.

References: 1. 11 B I.O.S. Final Rpt 933, Item 1 (1946), p A3/73
2. 11 B I.O.S. Final Rpt 933, Item 1 (1946), p A3/73
3. 11 B I.O.S. Final Rpt 933, Item 1 (1946), p A3/73

Ger 138

projectile was placed (below firing) over a spigot which was in the form of a short tube. At the base of the tube was inserted a cartridge case with a propellant and a primer. The firing was probably done in a manner similar to that for the Saiton Motor, i.e. by a striker held by a coiled spring and operated by a lanyard.

Spike Bomb. See Star-bomb.

Spillat (Spillat) A class of smokeless propellants prepared in 1898, by mixing sheets of paper and impregnating them with substances which slow down the rate of burning (wooden). The exact composition of these propellants was never revealed by the manufacturer, the Explosivfabrik Werke Spillat, Gieselschütz und Max Thom, Hamburg. The charges were made by superposing and compressing several sheets of untreated paper.

Reference: J. Daniel, Detonations, Paris (1902), p 735.

Spiltdwisch (Detonacy of Fragments). See Fragment Detonacy Test.

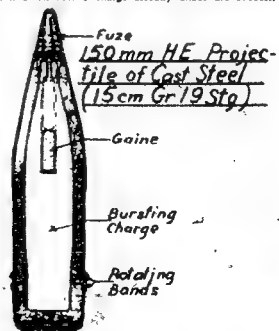
Splitting Process of Manufacture of Sulfuric Acid is briefly described under Sulfuric Acid Manufacture.

Sponting Powder. See Jagspulver.

Sponting Projectile (Schussrohrabzehrungsgranate). A projectile serving for observation and adjustment of artillery fire. It contained a small charge of smoke composition in a separate container inserted in the high explosive charge.

The following projectiles are described on pp 405, 494-5, 500, 529 & 535 of TM 9-1985-3 (1953):

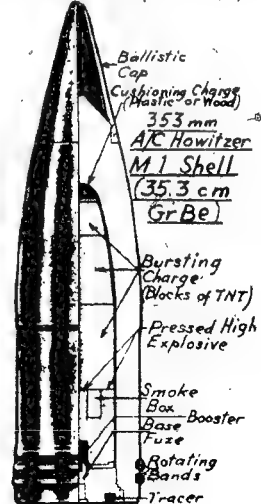
- 75 mm HE Projectile (7.5 cm Igr 18 AZ 23a) (or the Light Infantry Gun (IGG 18) or Light Mountain Infantry Gun (IGG 18)). It was about 13" long and contained 1.21 lb of an Anselm. Directly under the primer of the PETN booster (GZ24a C/98 Np) was located a small charge of smoke composition (pp 405-6)
- 150 mm HE Projectile 19 with Gaine 36 (15 cm Gr 19 mZdg 36) for Heavy Field Howitzer 18 (sFH 18). It contained 11.22 lb of cast TNT as a bursting charge and a small smoke charge directly under the booster.



Total weight of projectile was 95.7 lb. Two types of point detonating fuzes were used: AZ 23 or DoppZ 2/50. The base was provided with a screw-down plate (pp 494-5)

- 150 mm HE Projectile of Cast Steel (15 cm Gr 19 Stg) for Heavy Field Howitzer (sFH 18) and sFH (B) and for Heavy (Luft) Howitzer (sFH 18). It was similar in appearance to the previous projectile, except that it did not have the screw-down base plate (pp 494-5)
- 150 mm HE Projectile 19 (15 cm Gr 19) for Heavy Field Howitzer (sFH 18) and sFH (B) and for Heavy (Luft) Howitzer (sFH 18). It was about 29" long and contained 9.46 lb of TNT (in cardboard containers) as a bursting charge. A small charge of smoke composition was placed on the bottom of the shell. The projectile had a screw-down base plate. Two types of propellant fuzes were used: impact and composition (AZ 23 and DoppZ 2/50) and two types of boosters (GZ24a C/98 Np and GZ24a C/98) (pp 500-501)
- 353 mm Anzconcrete Projectile (35.3 cm GrB) for Howitzer M1 was conventional in design and contained 75 lb of TNT as a bursting charge and a small charge of a smoke composition used for spotting purposes. Total weight of loaded projectile was 1265 lb.

Note: According to information supplied by H.H. Bullock and A.B. Schilling of Picatinny Arsenal, it might be assumed that the HE filling consisted of four sections loaded in a cartridge. The 1st and 2nd front sections were cast TNT containing 54.07% was the 3rd section was cast anhydrous TNT and the 4th section was pressed TNT (or possibly



of sulfur recovery. (See under Lurgi Cracking Process.)

D. Jacob's Process was essentially as follows:

- Four vertical cylindrical packed extractors, fired with covers and each containing air tubes were loaded with spent oxides (7.5 tons in each vessel) and extracted with carbon disulfide at 25° entering each vessel at the top and moving by gravity.
- Of the 4 extractors, 3 were in the extraction cycle and one off for charging or discharging. As a freshly charged extractor was put on the line an extractor containing exhausted oxide was taken off.
- The freshly charged material was first treated with CS₂ rich in sulfur from there the saturated acid went to a 10 ft nozzle, water-heated still for distillation of fresh CS₂, from the head tank entered the most exhausted extractor.
- When the sulfur extraction in the spent oxide had succeeded in the economic limit, the extractor was taken out of the circuit and the CS₂ acid remaining removed by the still by direct injection of live steam at 5 atm pressure.
- After removal of the last traces of CS₂, the extractor cover was removed and the neck of stills lifted out.
- Distillation of CS₂ was conducted batchwise at 80-90° and the CS₂ was condensed and collected.
- When distillation was complete, the temperature in the still was raised to 130° by direct steam and the molten sulfur ran out through a jacketed pipe into a large shallow trench-way in the open air. Venting of the still was done with nitrogen.

A more detailed description of this process is given by H.A. Hoyle et al, BIOS Final Report 1644 (1948), pp 5-10.

Supergun. See Hochdruckpumpe.

Synonym. See Dinosebazine.

SW-Stell and Brenneise. According to CIOS Rept 10-115 (1945), p 11, the 90/10 mixture of concentrated nitric acid (Brenneise) to fumes made of ordinary nitric acid was used in conjunction with a combustible (Brenneise), such as gasoline, in liquid rocket propellants. The above acid mixture was known as SW-Stell. The same name was applied to the straight concentrated nitric acid (such as 98-100%) when used in rockets. This acid was also known as Selenic.

Synthetic Resins and Emulsions used in Germany during WW II for the manufacture of films employed in ammunition. The details are in BIOS Final Reports Nos 1715, 1794 and 1795 (1947).

Telfon. An experimental bilipid rocket designed to be fired in groups of 63 from a launching machine known as the Dehagen. The missile was about 21 in long and 3.8 cm in diameter. It was powered with a fuel containing 500 g of H₂. It was propelled by a liquid acid (Vical) and an oxidizer (concentrated nitric acid).

References:

1) CIOS Rept 29-36 (1946), pp 24-28

2) TNA 9-1945 (1935), p 223

Tapered Bone Gun (Wägrbüchse Gewehr), called also Tapered Bone Gun (Gewehr) or Reducing Bone Gun (Reduzierbüchse) was developed in the early stages of WW II. Its barrel consisted of 3 sections (starting from the breech):

- Cylindrical section, such as 42 mm bore diameter
- Slightly conical middle section and
- Cylindrical section, such as 28 mm bore diameter.

There were also guns with diameters 28 mm or 75 mm for section 2 and 20 mm or 55 mm for section 3.

Because of this construction, the projectile which

had a spoon-like body, was squeezed to a smaller diameter as it passed from the breech to the muzzle. The idea of this gun was to possess a larger cross-sectional area of the projectile to the propellant gases, and to possess a small cross-sectional area to the atmosphere in order to reduce air resistance and thus increase the muzzle velocity of the projectile. It was claimed that the most valuable advantage of this type of gun was the possibility of reducing the total length of a bore almost to one-half without any changes in maximum pressure and muzzle velocity and preserving almost the same weight of projectile.

Although this weapon was light and gave comparatively good underpenetration, it was given up for the following reasons:

- Its manufacture was very difficult
- It wore out too rapidly
- Its effective range was rather short.

Some of the tapered-bore guns and their projectiles are on display at the Aberdeen Proving Ground Museum, Maryland.

A short description of such guns is given by:

L.E. Supon, German Research in World War II.

J. Wiley, N.Y. (1947), p 189.

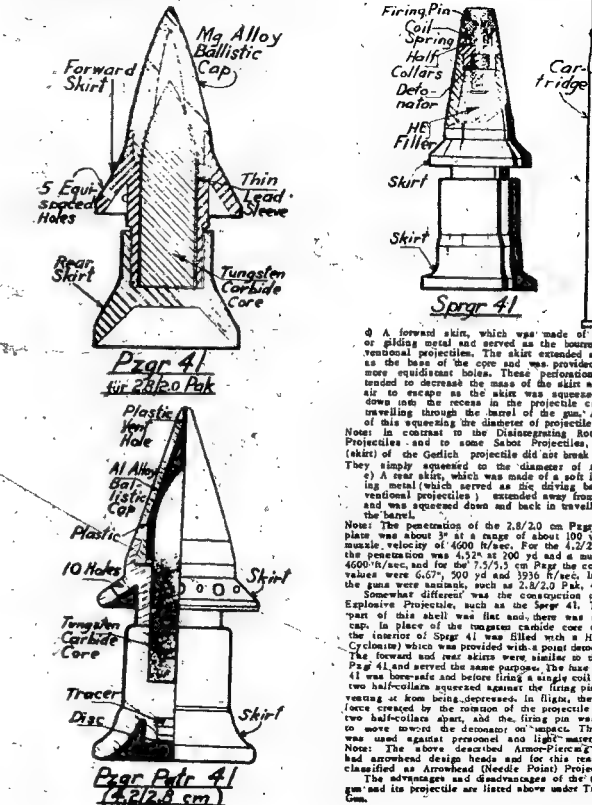
Note: According to E. Engelberg, The Ordnance Sergeant, May 1944, p 317, the inventor of this gun and its projectile was an American born German engineer, H. Gerlich, residing in Kiel. He worked on the development of high velocity weapons and projectiles from about 1920, and in 1932 he demonstrated at Aberdeen Proving Ground, Md a rifle firing a missile with a velocity of about 4445 ft/sec. The rifle was not accepted. After this Gerlich worked for the Germans. The first known combat use of the Gerlich principle was in the Libyan campaign. The weapon employed in Libya was the 2.8/2.0 Psk, a light antitank gun mounted on a recoilless carriage. In this gun the first 18° of the barrel, beginning from the breech, were of rather 2.8 mm, the next 30° of the barrel had a rapid taper of .022" per 1" and in the last 23° of the barrel, the taper decreased to .0022/1". The projectile had no grinding band or banded band, but instead had two slanting flanges extending away from the body. During the flight of the shell through the tapered bore, the skirts collapsed and a nearly smooth shell of about 20 mm caliber emerged from the muzzle of the gun. It was claimed that muzzle velocities up to 6000 ft/sec could be achieved and that armor penetration at 100 yds was 70 mm for hard steel and 76 mm (3") for machine gun plates.

Notes: According to TM 9-1895 (1935), p 360, the Sgweye "How Gun consisted of an ordinary rifled gun to the muzzle of which was attached a smooth-bore tapered extension. This meant that there was a difference between the Sgweye "How Gun and the Tapered Bone Gun. The projectiles were interchangeable in both cases. The guns and projectiles called "Sgweye-Bone" by the Americans were called "Telfon" by the British.

Tapered Bone Gun Projectile or Gerlich Projectile. According to E. Engelberg, Ordnance Sergeant, May 1944, pp 319-313 the typical Gerlich projectile such as the Arrowhead Projectile Type 41 (Psk 41) used in the 2.8/2.0 Antitank Gun (2.8/2.0 cm Psk) consisted of the following parts:

- A tungsten carbide core which had a diameter about half the caliber of the gun at the muzzle and served for the actual penetration into the armor.
- A thin lead sleeve which covered the core and held it in place. The sleeve served as a lubricant to the core when the skirts were separated from it on impact.
- A magnesium alloy ballistic cap which fired snugly into the forward skirt and served as the nose of the projectile. On impact the Mg alloy produced a flash which permitted observation of the firing.

Note: The Mg cap was not used in all tapered bore projectiles, as can be seen from the drawing of Psk 41. In this projectile the cap is of aluminum and the tracer composition, fitted into the base of projectile, permitted observation of the firing.



4. A forward skirt, which was made of a soft iron or gliding metal and served as the bourrelet of conventional projectile. The skirt extended as far back as the base of the core and was provided with 5 or more equidistant holes. These perforations were intended to decrease the mass of the skirt and to allow air to escape as the skirt was squeezed back and down into the recess in the projectile casing while travelling through the barrel of the gun. As a result of this squeezing the diameter of projectile decreased. Note: In contrast to the Disintegrating Hottel Sand Projectile and the Low Sabot Projectile, the bands (skirt) of the Gerlich projectile did not break on impact. They simply squeezed to the diameter of the muzzle.

5. A rear skirt, which was made of a soft iron or gliding metal (which served as skin diving band of conventional projectiles) extended away from the body and was squeezed down and back in travelling through the barrel.

Note: The penetration of the 2.8/2.0 cm Psk into armor plate was about 3" at a range of about 100 yards and a muzzle velocity of 4600 ft/sec. For the 4.2/2.8 cm Psk the penetration was 4.375" at 200 yd and a muzzle velocity of 4600 ft/sec, and for the 7.5/5.5 cm Psk the corresponding values were 6.625" at 200 yd and 1936 ft/sec. In all cases the guns were automatic, such as 2.8/2.0 Psk, 4.2/2.8 Psk. Somewhat different was the construction of the High Explosive Projectile, such as the Sgweye 41. The forward part of this shell was flat and there was no ballistic cap. In place of the tungsten carbide core of Psk 41, the interior of Sgweye 41 was filled with a HE (such as Cyclotol) which was provided with a point detonating fuse. The forward and rear skirts were similar to those of the Psk 41 and served the same purpose. The base of the Sgweye 41 was banded and before firing a single coil spring kept two half-collars against the firing pin, thus preventing at gun being depressed. In flight, the centrifugal force created by the rotation of the projectile forced the two half-collars apart, and the firing pin was then free to move inward the detonator on impact. The Sgweye 41 was used against personnel and light material targets. Note: The above described Arrowhead Projectile had arrowhead design heads and for this reason can be classified as Arrowhead (Nose Point) Projectile (v. 3).

The advantages and disadvantages of the tapered-bore gun and its projectile are listed above under Tapered Bone Gun.

The projectiles used in tapered-bore guns are also described in the following References:

- 1) R.A.D. Simon, *Pie Arm Tech Rep* 1326 (1944) (42/28 mm APBV).
- 2) A.B.Schilling, *Ibid*, 1378 (1945) (75/55 mm HE Shell for Tapered Bore Gun, Pat 41).
- 3) A.B.Schilling, *Ibid*, 1379 (1945) (75/55 mm AP Shell for Tapered Bore Gun, Pat 41).
- 4) *Dept of the Army Tech Manual*, TM 9-1985-5 (1953), pp 371-372: 28/20 mm HE, 28/20 mm, 42-28 mm HE and 42/28 mm AP projectiles.

Taken, see under Trinitro.

Target Indicating Flare, Mark, 50 (Rankin, and Target Indicator) are described in TM 9-1985-2 (1953), pp 71-74-84 (See also under Flare and under Marker).

Telluride (Tellurium). See under Cordite Charge Cartridge.

Television Guidance System for Missiles. See under Guidance Systems for Missiles.

Telluricopper oder Tellurische Mischschwamm (Plum Apparatus or Telluric Mixture Machine). An apparatus suitable for mixing solid and liquid ingredients of explosives, propellants and pyrotechnic compositions. It consisted of a large horizontal, cast iron, steam-jacketed, cylindrical pan on which the materials were placed. These were covered and mixed by the combined action of a long, small diameter, horizontal pillar (made from a non-sparking metal, such as Cu, brass, or Al) rotating around the center of the base at the rate of ca. 1 rpm and a series of scrapers (made from non-sparking metal) following behind the roller. The scraped material was removed by the roller and then again scraped and this action continued until all the ingredients were well mixed.

The apparatus was mounted before WW II by the Gehr, Burberg, Metzmann, and could be operated either in the cold, or heated by steam.

References: Sauerbacher, Schieser- und Sprengstoffe, Leipzig, (1955), pp 301-2.

Tellurium (Dislike Lead Mine). According to Simon (Ref 1) these mines gave the Allies considerable trouble throughout WW II. They were sufficiently powerful to "put a tank out of action and to wreck almost any other vehicle. The first of such A/T mines, called Tellurium 35, was made of steel, while the models developed towards the end of WW II were made of non-magnetic materials to render mine detectors ineffective. Some of the latest mines were reported to be remote-controlled but it is not known whether they were actually used in combat.

The following models are described in Ref 2: Tellurium 35 A/T (p 267); Tellurium 35 (Steel) A/T (p 268); Tellurium G A/T (p 269) and Tellurium-43, Pilot, A/T (p 270) (Pills mean mushroom).

Essentially the body of the mine was a circular, flat, disc-like form with a hole in the center of the cover. The body was loaded with 11-12 lb of compressed high explosive (such as TNT) and an igniter was screwed into the cover. A second (floating) cover was held down by a metal ring attached to the body and was supported in the center by a heavy spring. A pressure of 200-400 lb on the

the "floating" cover was sufficient to depress it as well as the igniter boning. The pressure of the boning on the top of the amber sheared the pin which held the striker in the cocked position, thus releasing the striker spring. As a result of this the striker set off the percussion cap, detonator, booster and the main charge such as TNT. References:

- 1) L.E. Simon, *German Research in WW II*, Wiley, N.Y. (1947), p 188.
- 2) *Amco, German Explosive Ordnance*, Dept of the Army, Tech Manual TM 9-1985-2, Washington, D.C. (1953), pp 267-70.

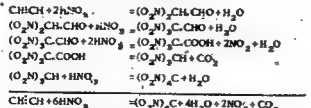
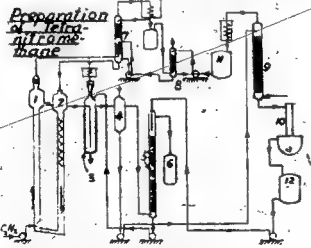
(See also under Landmine).

Testing Gallery (Schlagversuchschachtele). See general section, under Galleries; Testing and also this section under Versuchsachtele.

Tellur oder X-Sellf (Tellurmonothane, abbreviated in this work as TeMe or TeMeMe). A detailed description of the preparation, properties and uses of TeMe is given in the general section under Methane. The following description concerns the German method of preparation and uses of TeMe.

As the classical method of prep. of TeMe from acetic anhydride and nitric acid (see general section) is very expensive, a new method was developed during WW II by Dr Schimmelmann (Refs 1 & 2). The laboratory scale procedure was as follows:

In an all-glass apparatus, schematically represented in the enclosed drawing, acetic acid reacted with nitric acid to give nitroform and the mixture of nitroform and nitric acid yielded TeMe on heating with sulfuric acid. The reaction being intended to proceed as follows:



About 60% of acetylurea reacted as above and about 40% underwent complete oxidation according to the equation $\text{CH}_3\text{COOH} + 2\text{HNO}_3 \rightarrow 2\text{CO}_2 + 2\text{H}_2\text{O} + \text{H}_2\text{O}$, so that the over-

all equation could be represented by:



The recovery of nitroform and of unconverted nitric acid was about 96% of theory.

In the procedure the acetylurea gas, $\text{C}_2\text{H}_5\text{O}$, was introduced at the lowest point of the system at the rate of 95.5 liters per hour and the nitric acid (98%) containing mercuric nitrate as a catalyst, was fed at the rate of 2.4 liters per hour.

Note: The catalyst was prepared by dissolving 70 g of mercury in about 100 ml of 98% nitric acid, adding 300 ml of water and making up to 1 liter with 60% nitric acid. Twenty ml of this solution was added to every 10 liters of 98% nitric acid.

By circulating cold water through the cooling coil, located in the 2nd leg of the reaction system, a temperature of 50° was maintained. The solution of nitroform in nitric acid overflowed from the circulating system to give nitroform vessels (3) placed in series, each containing a stirrer heated by a steam jacket. The sulfuric acid from the TeMe pump (5) together with the nitroform mixture flowed through the last section of the steam-heated vessels and the remaining acid overflowed from the 1st nitroform vessel.

In each section of the system, the reaction mixture was maintained at 50° and the top layer of TeMe was continuously removed. The reaction mixture was provided with a reflux condenser for returning TeMe and HNO_3 as well as any condensible gases such as N_2O .

The waste mixture leaving the 3rd nitroform vessel separated in (4) and the top layer of TeMe was fed continuously to the purification column (5). The feed of 95.5% sulfuric acid to the purification column (5) was 1.7 liter per hour and the product was charged to the strikers (6) and (7). Pure TeMe left the top of the purification column (5) at the rate of 440-460 g per hour and was collected in a tank (8).

The off-gases of the nitration system (such as N_2O , CO_2 with small amounts of HNO_3 , CHNO_2 , CHNO , CH_2O and possibly unreacted $\text{C}_2\text{H}_5\text{O}$) passed to the purification column (7) which was divided into 2 sections. In the lower section the last traces of $\text{C}_2\text{H}_5\text{O}$ were removed by scrubbing with warm nitric acid (containing mercuric nitrate) fed at the rate of 2.4 liter per hour. In the upper section of column (7) nitrogen dioxide and carbon dioxide were separated by distillation and the nitrogen oxide was condensed.

The gases leaving the receiver were scrubbed in a smaller column (8) by cold nitric acid (to remove the last traces of nitrogen dioxide) and the nitric acid run-off was fed to column (7), whereas the CO_2 was allowed to escape.

The nitric acid (which contained sulfuric acid, nitrogen dioxide and tellurmonothane) was separated from sulfuric acid by distillation in column (9) and condensed in tank (11). The residue (consisting of 70% sulfuric acid) was concentrated to 95.5% in column (10) and collected in tank (12).

Note: Although the attached diagram indicates a continuous system for the preparation of TeMe, the process was actually conducted batchwise, as sufficient material accumulated, TeMe was produced at a later very powerful and TeMe explosive, called in Germany *Tellurmonothan* (A/T), and also as a very powerful and TeMe explosive, called in Germany *Tellurmonothan* (A/T), and also as a very powerful and TeMe explosive, called in Germany *Tellurmonothan* (A/T).

It was proposed to use this mixture in V-Z rockets (Ref 2).

References:

- 1) R.E. Schimmelmann et al., *COS Rep* 25-48 (1945), pp 6-14.
- 2) *Wissenschaftl. u. Techn. Fortsch. Rep* 7 (1946), pp 1-2.

Tellur Sprengstoffe (Tellurmonothane Explosives). It was employed in the German mortar shells, but in the acute shortage of TNT and of other high explosives, the German used during WW II, as ingredients of explosive mixtures,

substances which were not explosives. Among such substances was TeMe (tellurmonothane), called in Germany *Tellur*, a liquid waste product of very finely pulverized aluminum (called *Pyroschiff*), impregnated with TeMe, and a small amount of the following substances:

a) a hydrogenation resin in hydrogen and a consolidating compound called *K₂*, which was a high dispersion of silica prepared by a special process. The hydrogenation was added in order to increase the sensitivity to initiation. This *Tellur* explosive was a solid possessing a very high

explosives with such properties were found to be suitable for underwater explosions (Ref 1).

Other explosive mixtures consisted of *Tellur* with liquid fulminating substances, such as hydrazine, coal, charcoal, etc. Some of these mixtures were more powerful and resistant than TNT, PETN, RDX, and were particularly suitable for underwater explosions.

Considerable work on this subject was done by Dr A. Schimmelmann (See general section under Methods). One of the most powerful and resistant explosives, known is a mixture of *Tellur* with fulminic acid, its velocity of detonation is about 9300 m/sec.

There were also explosives prepared from derivatives of TeMe, eg for instance the perchloric ester of trinitroethane. The trinitroethane was prepared by condensing nitroform (derived from TeMe) with 40% solution of formaldehyde.

References:

- 1) G.E. Jones, *PBL Report* 85-160 (1945), pp 2-3.
- 2) A. Schimmelmann, *Schieser- und Sprengstoffe*, Leipzig, (1955), p 185 and 186; Spreng- und Schmelzmittel, Zürich (1948), pp 10, 16 & 148.

Tetramine (Tetrazin) was used in Germany utilizing the same equipment as used for prep. of L.A. and L.Sz.

The procedure was as follows:

- a) To a solution containing 40 kg of Na nitrate in 150 liters of hot water, 60 liters of water were added to 50° and was added gradually and with stirring 40 liters of an aqueous solution of 5.9 kg of aminopuridine sulfate (the addition rate was 1 kg per hour).

- b) After stirring the mixture for an additional hour at 50° and for 1 hour at 20°, the reactor was tipped and the contents caught on a filter cloth made of horse hair.

- c) After washing, the ppt with several portions of water, it was dried in the same manner as described under lead azide. This gave about 3.0 kg of dry tetrazine.

- d) Boiling the mother liquor for several hours was sufficient to destroy any warm tetrazine remaining in it.

A similar method, used at the Fabrik Wolfenbutten Chemische Werke (and at the Stadel, Berlin, Dynamit A.G.), is described by Sheldon (Ref 3). In this description the following details of the method which are worthy of mention are given:

A. A solution of aminopuridine sulfate (5 kg per 40 l of water) was neutralized (to the litmus paper end point) with sulfuric acid or nitric acid and then added to a prepared solution of Na nitrate (2.5 kg per 50 l of water). If the addition rate was rapid, slow stirring crystals were formed. If the addition rate was slow (2 hours), larger and fewer crystals resulted.

B. The detailed procedure as follows: (temperature of 30 to 35° was maintained throughout the entire reaction period which was allowed to proceed 30 minutes after the feed of the aminopuridine sulfate solution had been added to the reactor. Then the agitator was stopped, the product allowed to settle and the mother liquor removed

Trinitrophenol (Trinitrophenol) (TNP), and Dininitrophenol (Dinitrophenol) (DNP) were used by the Germans during WW II in some composite explosives. They were manufactured at Semina Fabrik, Pardubice, Czechoslovakia. See also general section under Naphthalene.

Trinitrorescin (Trinitrorescinol) (TNR), or Syphalic Acid. See Trinitro.

TRINITROTOLUOL oder TROTYL (Trinitrobenzol) (TNT) $\text{C}_6\text{H}_2(\text{NO}_2)_3$ oder Fo O3 (Filler 1902, $\text{C}_6\text{H}_2(\text{NO}_2)_3$) is described below fully in the general section under Toluenes.

TNT was officially adopted in Germany in 1902 as a military explosive, earlier than in any other country. Its aerial use by the Army was begun in 1904, and the industrial production started in 1906 at the Schleichbach Fabrik, D A G.

For the description of German methods of preparation of TNT, as practiced before, during and after WW II, see the books of Reichel (Ref. 1) and Schleichbach (Ref. 2). The same books give also the properties of TNT.

It is to be noted that before and during WW II, the Germans used a rather complicated process for the manufacture of TNT. This was due to the fact that tolerance in these days was rather impure. This method, described by Reichel (Ref. 1, p. 377) was briefly as follows:

After dissolving toluene by means of weak nitric acid-sulfuric acid to produce MNT (mononitrotoluene), the crude product (monooil) was separated from the monosulfuric acid, then washed with water and finally with weak sodium hydroxide solution. After blowing the live steam through the oil (in order to remove the benzene present) as an impurity as well as any unreacted nitric acid, it was cooled to allow the p-MNT to crystallize (a p. 378). After separating the p-MNT by filtration, the remaining liquid fraction was subjected to fractional distillation under vacuum using a column apparatus. The o-MNT came off first, leaving the m-MNT as a residue. Only p- and m-MNT were used for the purpose of military grade TNT. The o-MNT was used for the purpose of liquid-DNT-TNT mixture (Dip oil) useful as an ingredient of commercial explosives. Another method to distill the o-MNT from the washed monooil and then to add the residue in order to separate the p-MNT from m-MNT.

The method of purification of TNT proposed by the Chemische Fabrik Gries was described in Ger P 207-170 (1908).

During WW II the German capacity was as much as 55 million pounds of TNT per month, but the maximum they ever produced was 49.5 million in April 1944. The TNT used by the German Army had a p of 80.4-80.5.

The manufacture of TNT during WW II in various German plants is described by Sickland et al. (Ref. 3 and 4) and Brooks (Ref. 6, pp. 39-41). It seems that some of the processes used in Germany was an efficient (from the point of view of speed of manufacture and yield) as the process introduced during WW II into this country at Keystone Ordnance Works, Meadville, Penna., by Dr L. A. Genoff, and finally adopted by all U.S. Ordnance plants. The maximum German yield was about 700 tons TNT per 100 p of m-Nitro, while the American yield was as high as 210 p (average yield was 205-208 p).

In one of the largest German plants, the Krimmel Fabrik of D A G, the following batch method was used during WW II:

- Pre-nitration.** The mononitro acid (consisting of 28% HNO₃, 56% H₂SO₄, and 16% H₂O) was added to the charge of toluene in the reactor 2.5 parts of acid to 1 part of toluene. The temperature was maintained at 35-40° by cooling coils and a jacket.
- Post-nitration.** The mixture was transferred to a post stirrer where it remained for several hours at 35-40°. Total time required for a full charge of m-Nitro (to be) was 5-6 hours.
- Separation.** The mixture was transferred to a cast iron vessel where it was allowed to stand for 6 hours. The mononitro acid (N₂O₄ 0.5%, H₂SO₄ 70% and a small amount of nitric acid) was separated and went to the acid recovery plant while the oil underwent purification.
- Purification.** The crude oil was washed with water until nearly neutral and was then steam distilled in the presence of NaOH (1% NaOH based on the total weight of m-Nitro). The purpose of adding NaOH was not only to neutralize the remaining traces of acidity but also to transform the nitrobenzene, present as impurities, into sodium nitrobenzenes, which are soluble in water. During the distillation the first fractions were collected separately because they contained some unreacted toluene, benzene, and other volatiles. After separating the m-Nitro from the water-soluble fractions, it was through caustic soda washes where the last traces of nitric acid were removed to nitrobenzene. The damp around m-Nitro (yield 138-145%) was forced by compressed air through large tanks to be ready for distillation.
- Distillation.** The product separated from impurities consisted of 96% o- and p-MNT and 4% m-MNT. The purification procedure took about 2 hours. Total time for the preparation of the m-Nitro was 13-14 hours, which was much longer than the present American practice.

B) Distillation or Distillation consisted of the following steps:

- Pre-nitration.** A charge of m-Nitro was mixed with bi-sulfuric acid (previously diluted slightly with water to separate the excess part of dissolved DNT) in order to use up any residual HNO₃ as well as to extract the last traces of DNT.
- Mixing.** After separation from the dilute acid, the m-Nitro was fed into the distiller containing the tri-sulfuric acid, consisting of 45% HNO₃, 3-4% N₂O₄, and 80% H₂SO₄, and cooled to 30°. During the addition of the m-Nitro the temperature rose to 60-65° and then fell to 55° due to the excess of unreacted m-Nitro.
- Post-nitration.** In order to complete the nitration, 60-70% nitric acid was added to the above mixture, and the temperature was allowed to rise to 70-72°.

Note: Time required for total distillation was not given. In order to ascertain if the mixture was completed, a sample of distillate was taken and distilled plate steam. If so, m-Nitro distilled off, the mixture was considered complete.
- Separation.** After allowing the charge to stand for 1 hour, the oil was pumped and transferred to an intermediate storage vessel, while the distillate (ca. 5% N₂O₄, 0.6% HNO₃, 78-80% H₂SO₄) was slightly

diluted with water in order to separate the greater part of the DNT and to obtain an acid consisting about 4.5% N₂O₄, 0.5% HNO₃, and 73% H₂SO₄. This diluted acid was mixed with m-Nitro, as was mentioned under (a). After this, it was transferred to a storage tank where it was allowed to remain for 4-5 days before being sent to the acid recovery plant. Some additional oil separated out during the storage. Note: Distillation in the recovery house of the distiller, as well as of the mono-waste acid mentioned previously, gave weak nitric acid (50-55% HNO₃) and weak sulfuric (68-70% H₂SO₄).

C) Trinitration. In the older Krimmel plant, the acid was added to the oil while in the newer plant the reverse procedure was used while the current American practice. The new method was essentially as follows:

- Mixing.** The trinitration was charged with oximized acid (HNO₃ 24%, H₂SO₄ 78%) at a temp. of 74-76° and the oil was added gradually, with agitation; while the temp was allowed to rise to 84-85°. The reaction was completed by raising the temp to 90° and maintaining it there for about 4 hours. Total time of nitration was about 6 hours.
- Separation.** The agitation was stopped and the mixture allowed to settle for ½ hour. The oil containing residual acid (1-2% HNO₃ and 1-2% H₂SO₄) was transferred to a washing house and the prepurified acid was slightly diluted with water (in order to precipitate out some additional TNT) and this diluted acid was used for the denitration (see above).

Note: Each nitration house was provided with an individual fume recovery plant. The gases formed in the nitration were removed through ventilators and forced into absorption towers where they were sprayed with water, thus forming weak nitric acid (50-55% concentration). This acid was removed for use in the mononitration.

D) Purification of TNT consisted of the following operations:

The oil (called Rohol) was given several washes at 90° and then neutralized at 80° with bi-carbonate of soda. The resulting product had a setting point of 78-78.5°. The melting product had a setting point of 78-78.5°, much lower than for pure TNT (80.4°) due mostly to the presence of unreacted m-Nitro, DNT, and other impurities. For further purification, the neutral Rohol was stirred with an equal amount of water at above 80° and the emulsion cooled to 74-76° with constant stirring to effect crystallization. At this point a saturated solution of Na sulfate (Sulfate) was added with continuous stirring. The resulting slurry was filtered and the precipitate washed with water.

Note: The Sulfate treatment removed the isomers of TNT (mostly beta- and gamma-isomers) to the amount of 4-4.5%, tetraminotoluene (TetM) present to the amount of 0.2-0.3% and some other impurities. Total loss from this treatment was 6 to 8%. The resulting product, called Rohol, had a setting point (a p) between 80.0 and 80.5.

E) Drying, Picking and Packing operations were conducted as follows: The purified TNT was heated to 85-90°, separated (while in the molten state) from water and then dried in special water-based vessels by bubbling dry air (at 85-90°) through the molten mass. The molten TNT could be sent from the driers either

directly to a shell-loading plant or to a flaker. The product with a p. of 80.5 or higher was called Grade A, that with a lower p. was Grade B. There was also a Grade UX (uncombustible) with a p. of 80.7-80.8° which was prep by recrystallizing Grade A TNT from a water emulsion, recaking the crystals with a small amount of sulfate, rinsing with water and drying.

The yield at the Krimmel Fabrik was 138-142 parts of pure TNT per 100 p of m-Nitro, or 200 p TNT from 100 p of toluene.

Capacity of the Krimmel Fabrik was 3,000 metric tons per month.

Brooks (Ref. 6) and Wendes & Lisle (Ref. 10) describe the following method of manufacture of TNT at the Allied Chemical Co. of Dynamit A G:

The mononitration method consisted of the following:

- Mononitration (continuous process)** was conducted in two stages. Toluene and nitric acid were fed into two pre-nitrators where the mixture was vigorously agitated for ½ hour at 35°. About 93% of the nitration was accomplished in this reaction. Toluene was fed in at a rate of 1,000 lb per hour. The resulting emulsion overflowed into one main nitration and then to a continuous gravity separator, where a recuperative gas packed with Raschig rings. The mononitro-waste acid was drawn off through a trapped bottom outlet while the mononitro went to a washer. Here the oil was washed with water and soda-ash solution and then passed through a series of stripping towers. Live steam was blown through the towers to remove the impurities, such as unreacted toluene, benzene and paraffins. The refined mononitro was then sent to the bi-nitration or shipped to other TNT plants.
- Bi- and Trinitration (batch process)** was conducted in much larger agitators than used in the U.S.A. As much as 10,000 lb of mononitrobenzene was treated in one batch (about 3 times as much as in the U.S.A.).

The bi-nitration took about 3 hours while the trinitration required 6 hours. For this reason there were twice as many bi-nitrators as bi-trinitrators. In the trinitration, the mixed acid (consisting of nitric acid 24%, sulfuric acid 78% and water 9%) was added to the crude DNT (Rohol) while maintaining the temperature at 83°. Then the temperature was raised over a period of 10 minutes to 98° and maintained at this point for 2 hours.

Note: There were no bottom outlets in the nitration, permitting the draining of the charges, but in case of fire there was a quick-opening valve which permitted large streams of 95% sulfuric acid to spray into the agitator to extinguish the fire (Ref. 6, pp. 9-10).

C) Purification of TNT (at Allied Chemical). Trinitro was washed with hot water, and then crystallized from fresh hot water. After drawing off the water and recrystallizing the product, it was treated with a 14% solution of Na sulfate (Sulfate) of pH 5 to 6 in such a quantity that there was from 3 to 4 lb of Na sulfate per each 100 lb of TNT. When the 14% soln was mixed with the TNT slurry, there was sufficient water present to keep the strength of solids to about 3% of Na sulfate. The red water was filtered off leaving a TNT with setting point 79.5 to 80°. For a pure product (a p. about 80.5°) the partially purified TNT was reinitiated

von Torpedos, Mines, und Tiefenbomben unter Berück-

WEAPONS (Weapons) may be subdivided into:

- A. Small Arms (Handfeuerwaffen), which include:
 - 1. Pistol (Pistole) revolver (Revolvier), carbine (Karabiner), rifle (Gewehr), machine gun (Maschinengewehr), and sub-machine gun (Maschinengewehr) models.
 - 2. Artillery Pieces (Geschütze), which include:
 - a. Cannon (Kanone), mortar (Mörser) and mortar (Mörser) models.

C. Rocket Launchers (Raketenschusswaffen), which include:

- 1. Panzerfaust, Panzerfaust, Panzerfaust (Raketensprengwaffe), Püppchen (Raketensprengwaffe) and others.
- 2. Most of the German weapons used in WW I and II may be found on display in the Museum of Aberdeen Proving Ground, Maryland.

Table 61, following, gives some of the characteristics of German small arms, artillery pieces and rocket launchers.

Table 61 (Weapons)

Caliber and Designation

6.35 mm (250) Mauser Automatic Pistol M 1910, called Maschinepistole (M.P.), Vest Pocket Pistol

6.35 mm Walther Pistol Models 1 (1908) and 2 (1910)

6.35 mm Walther Pistol Models 5 (1913), 8 (1920) and 9 (1921)

6.5 mm Pistol: Bergmann, Original, Sauer and others

6.5 mm (250) Bergmann Automatic Pistol

6.5 mm Mauser Vest Pocket Automatic Pistol, Types WTP (1910) and WTP II (1919)

6.5 mm Sauer & Sohn Vest Pocket Pistol, Types I and II

7.65 mm (300) Military Mauser Automatic Pistol, called Maschinepistole (M.P.) developed in 1895 and used during WW I

Note: According to Ref 8, 41, p 177 there was also an improved model (M1926) of the above pistol

7.65 mm Mauser Machine Pistol M 1932, called Schmelz-Pistol (Rapid-Fire Pistol) issued to SS troops. Was also issued in Spain under the name of ASTRA

7.65 mm (301) Automatic Pistol, introduced in Germany in 1893 by an American, Hugo Borchardt

7.65 mm Mauserlicher Pistol invented in 1900

7.65 mm Luger (Parabellum) Pistol M 1900 and M 1900/06 were used during WW I. Model 1900 was an official Swiss pistol

Note: According to Smith (Ref 9, p 462) the original Luger was designed by an American, Borchardt, and was further developed by a German, Luger. It was first issued under the name of "Borchardt-Luger" and then changed and shortened in the U.S.A. to the name "Luger". The name "Parabellum", which literally means in Latin "for war", is used in Europe. See also p 9

7.65 mm Automatic Carbine (Parabellum Karabiner)

7.65 mm Dreyse Automatic Pistol M 1907

7.65 mm Bohalla Automatic Pistol would be Becker & Heilbrunn, Sol

7.65 mm DWM Automatic Pistol, named by the Deutsche Waffen- und Munitionsfabrikanten

7.65 mm Automatic Pistol (invented by F.L. Schmidt of Sol and called F.L. Schmidt's (F.L. Schmidt's)

7.65 mm Automatic Pistol, called PB Special Model II, named by A. Meiss, Sol

7.65 mm Original Automatic Pistol (named by the Deutsche Waffen- und Munitionsfabrikanten)

7.65 mm Jäger Automatic Pistol

7.65 mm Mauser Automatic Pocket Pistol M 1910

Remarks, Uses and Some Characteristics

Length: barrel 2.0 ft and overall 4.0 ft; w 10.22 oz and capacity of 6 rounds. One of the low small pistols ever produced

Blackjack vest pocket pistol using .25 CAC

Streamlined versions of above pistols

Can be seen in the Museum of Aberdeen Proving Ground, Md.

One of the earliest small size pistols

Elementary blackjack pistol resembling the Browning type. The Type II was the streamlined version of Type I

Resembled a Browning in external appearance. Capacity 7

Recall-operated pistol weighing 45 oz. Capacity 10. Could be fired with shoulder stock holster attached

Was also an improved model (M1926) of the above pistol

Recall-operated weapon which may be considered as intermediate between the pistol and the sub-machine gun. Length of barrel 5 1/2 ft, overall 12 ft, w 45 oz, capacity 10 or 12 cartridges, as fed up to 1500 ft/sec

Considered as the forerunner of the Luger. Could use 7.65 mm Mauser ammunition

Was also made in caliber 7.65 mm

Barrel length 4 1/2 ft. Used cartridges cost 10 pf. w 27 in 22 M and 1 bullet weighing 93 gr. Mx vel 1250 ft/sec

It consisted of a regular Luger pistol provided with a detachable wood stock and a long barrel with a checkered wood grip

Blackjack-type pistol weighing 24 oz. capacity 8

Blackjack-type pistol weighing ca 22 oz. For hand carrying with WTP design

Blackjack-type pistol weighing 20 1/2 oz. Capacity 7

Blackjack-type pistol weighing 22.0 oz. F.L. Schmidt of Sol and called F.L. Schmidt's (F.L. Schmidt's)

Double-action blackjack pistol which closely resembled Walther's DPK

Stoker-fired blackjack pistol

A blackjack-operated pistol of simple and most unusual design

A straight blackjack-action pistol weighing 21 1/2 oz. Capacity 4

References

2, p 321; 4, pp 275-8 & 10, v1, pp 141 & 560

11, p 478

11, p 478 & Ref 12

7, p 27

11, p 485

11, p 484

2, p 321; 4, pp 275-8 & 10, v1, pp 141 & 560

8, v1, p 177 & 11, pp 468-71

7, p 27 & 10, v1, p 185

7, p 27

2, p 320; 3, p 187; 7, p 27 & 10, v1, p 185

10, v1, p 184

10, v1, pp 235-5 & 282 & Ref 12

10, v1, pp 218 & 379

10, v1, pp 235-6

10, v1, pp 245-5 & 385

10, v1, pp 253-4 & 388

10, v1, pp 254-6

10, v1, pp 249-3 & 385 & Ref 12

10, v1, pp 246-9 & 387

WEAPONS (PISTOLS AND REVOLVERS)



PICATINNY ARSENAL
TECHNICAL REPORT NO. 2510

DICTIONARY OF EXPLOSIVES, AMMUNITION AND WEAPONS

(GERMAN SECTION)

BASIL T. FEDOROFF

HENRY A. AARONSON GEORGE D. CLIFT EARL F. REESE

DOVER, NEW JERSEY

1958

Ger 229

(Weapons) (Cont'd)

Remarks, Uses and Some Characteristics

Double-throw blowback pistol, length barrel 3.98" and overall 6.5"; Wt 20.6 oz and capacity 8 cartridges, either 7.65 mm Browning or .32 CAP

High-capacity pistol weighing 23.6 oz. Long recoil-operated weapon, weight 23 oz with capacity 7 cartridges, caliber .301

Was replaced after WW I by M 1930 and M 1938

Blowback-operated weapons, capacity 7. The Bekendtschuss was widely used by military and police officials

Streamlined modification of earlier models

Slight blowback-action weapons. Length of barrel 3.5" and overall 6.5", Wt 22 oz, capacity 8 cartridges either 7.65 mm Browning or .32 CAP

Blowback-action weapons using .32 CAP cartridges

Valuet pistol widely used by police forces throughout Europe

Designed for detectives who carry this weapon concealed

Prototype of Army rifle used in both WWs. The first 500,000 rifles were made in 1888 by L. Krupp & Co., Berlin. The carbine (Karabiner) was slightly shorter and lighter than the rifle. Both of them used rimmed, pointed, center-fire cartridges with mild steel bullets

Length of barrel 29.13" and overall (without bayonet) 49.25"; Wt 9.5 lb. Capacity 5 daisy-bellows-magazine cartridges with pointed bullet (Spitzer). Muzzle velocity 2807 ft/sec and pressure 3500 atm (51.13 psi)

Length of barrel 29.13" and overall (without bayonet) 49.25"; Wt 9.5 lb. Capacity 5 daisy-bellows-magazine cartridges with pointed bullet (Spitzer). Muzzle velocity 2807 ft/sec and pressure 3500 atm (51.13 psi)

Cavalry version of Gew 98. Barrel length 18"

Carbine version of Gew 98. Length of barrel 24" and overall 43.5"; Wt 8.2 lb. Capacity 5

Slightly modified version of Kar 98. Was used in WW I

It differed from Kar 98 in having a hinged bolt handle and side sling. Was used during WW I

Can be seen at the Museum of Aberdeen Proving Ground, Md

Can be seen at the Museum of Aberdeen Proving Ground, Md

Slightly modified version of Gew 98 designed to permit operating by means of reducing machine operations

Was provided with magazines of 10, 15 and 25 round capacity

Short recoil-operated, water-cooled MG used during WW I. Wt 45.5 lb with feed

A lighter version of MG 08, which weighed 30 and 31 lb. Its air-cooled version, made at Spandau Arsenal, was called Spandau Machine Gun

References

10, v.1, pp 246-9 & 247 & 11, pp 472-3

10, v.1, pp 246-6

10, v.1, pp 258, 260 & 11, pp 483

10, v.1, p 259

10, v.1, pp 279-8, 282-4 and 11, pp 476-7

10, v.1, pp 286-7 & 11, p 478

10, v.1, pp 286-7 & 11, p 478

10, v.1, pp 286-92, 32; p 478 & Ref 12

10, v.2, pp 201-13, 11, pp 475-7 and Ref 10

4, pp 83-90, 10, v.2, pp 171 & 11, p 429

10, v.2, pp 171 & 11, p 429

11, p 428

4, pp 85-90, 10, v.2, pp 171-75; and 11, p 428

10, v.2, pp 171 & 11, p 429

10, v.2, pp 171 & 11, p 429

12

12

10, v.2, pp 175-6

10, v.2, pp 176-7

4, v.1, pp 309 & 11, p 462

4, v.1, pp 309 & 11: 11, pp 472-20 and Ref 12

Ger 230

WEAPONS

(CARBINES AND RIFLES)



Caliber and Designation

7.65 mm Mauser Automatic Pistol, M5c (Mauser-Caliber)

7.65 mm Rheinmetall Automatic Pistol

7.65 mm Roth-Sauer Automatic Pistol was somewhat similar to the Luger-Huganau 200-Steyr pistol

7.65 mm Sauer Automatic Pistol M 1908 made by J. Sauer & Sohn, Suhl

7.65 mm Sauer Automatic Pistol M 1913 and Behrendtschuss (Austrian Model)

7.65 mm Sauer Automatic Pistol M 1930

7.65 mm Sauer Double Action Automatic Pistol M 1930 (called star Model 19 was widely used during WW I by the German air mail tank forces. Considered one of the world's best pocket pistols

7.65 mm Walther Pistol Model 3 (1909), (GROD), Grand 7 (1917), made by K. Walther of Zeitz, Meissen

7.65 mm Walther Pistol PP (Polizei Pistol), introduced in 1921

7.65 mm Walther Pistol PPK (Polizei Pistole Kriminal) introduced in 1929 and made in great numbers

7.9 mm (.311") Rifle M 1888 (Gewehr 88, abbreviated to Gew 88) and developed by a German Military Commission. It combined a modified Mauser (M1871) receiver bolt system with a modified Mauser loading system (magazine)

7.92 mm (.312") Mauser Rifle M1898 (Gewehr 98), Bolt Action, was the standard German infantry rifle of WW I and the ready form of all modern Mauser rifles. Served as prototype for military rifles of many European and South American countries

Mauser Original Gew 98 had a round nose bullet (some as in M 1898 which had a slightly smaller than the pointed bullet. In order to make the new bullet it was necessary to enlarge the die of Gew 98

7.92 mm Mauser Carbine 1896 (Karabiner 96, able to Kar 98). Original model

7.92 mm Mauser Carbine 1908 which was introduced in 1904 and adopted in 1908 for use by military and civilian (sporter) personnel

7.92 mm Kar-98 was introduced after WW I by the Reichswehr

7.92 mm Kar-98, developed after WW I by the Reichswehr for cavalry and mounted forces use

7.92 mm Karabiner 98 h (Kb-98h), Model 1913

7.92 mm Semi-Automatic Rifle, Model 1913

7.92 mm Gewehr 98/17, developed during WW I and discarded after it

7.92 mm Gewehr 18, developed after WW I as an experimental model

7.92 mm Machine Gun M 1908 (MG-08)

7.92 mm Machine Gun M 1908/15 (MG-08/15) Maxima

Caliber and Designation

7.92 mm Gewehr 13/40 (Modification of Czech Model 39)

7.92 mm Gewehr 98/40 and 79/40 Mauser

7.92 mm Amstutz Rifle, Model 55-41

7.92 mm Semi-Automatic Rifle Model 41-41 (Haitzschmatische Gewehr 41-41) developed at Mauser plant

7.92 mm Semi-Automatic Rifle Gew-41 (G-41) and its improved version G-41W were designed by

7.92 mm Semi-Automatic Rifle M 1943 (Gew-43) and Carbine M 1943 (Gew-43) were developed during WW II in order to do away with some defects of G-41 and G-41W weapons

7.92 mm Automatic Rifle, M 1942 (Light Machine Gun), called "Schmitz" or Gewehr 42 (Panzertruppen Rifle 42), able to FG42. It was fitted with a folding bipod mount

Note: This weapon was made by the H.K. Schmitz Waffenfabrik, Suhl. It was also made in the U.S.A. under the designation of T-44

7.92 mm Automatic Rifle, M 1942, Modified

7.92 mm Light Machine Gun, MG-42 was the latest German machine weapons of WW II and the most remarkable gun of its type ever produced in any country of the world.

MG-42 incorporated the best features of previous Russian and German MGs

7.92 mm Machine Carbine (Machinecarabine), able to MG-42

7.92 mm Machine Carbine MKb-42 (H) and MKb-42(W), called also Submachine Gun

7.92 mm Carbine 43, Kb-43

7.92 mm Machine Pistol M 1944 (Maschinenpistole 40), was originally developed in 1942 and then improved in 1944. On Hitler's order it was called "Sturmgewehr 44 (StG-44)

Note: The cartridge used in the latest 7.92 mm weapons, such as machine carbines and machine pistols, was a cut-down version of the standard bottle-neck rifle cartridge using a 125 grain pointed bullet. Muzzle velocity was ca 2750 ft/sec ca 550 yd (Ref 11, p 502)

7.92 mm People's Rifle 1 (Volksturm Gewehr 1, able to VG-1), made by K. Walther, Suhl

7.92 mm People's Rifle Special (Short) was developed in 1942 by K. Walther and introduced in 1942

8 mm (.311") Schwarzenegger Machine Gun M 1907/12, invented by A. W. Schwarzenegger of Germany and first made by the Steyr Arms Works in Austria

Remarks, Uses and Some Characteristics

Short weapon (barrel 18") used by mountain and ski troops

Can be seen at the Museum of Aberdeen Proving Ground, Md

Same as above

Gas-operated weapon which did not prove to be successful in field use

Experimental gas-operating weapons incorporating some features found in pre-WW II Russian Design, Simonov and Tokarev weapons

These weapons were gas operated and the action was of the straight-line (non-rotating) bolt type. Characteristics of Gew 43: overall length 44.5", barrel 22", wt 9.9 and magazine capacity 10 cartridges from two Mauser weapons

Gas-operated, air-cooled weapon of revolutionary design. Overall length (without bayonet) ca 42", barrel ca 19" and wt 9.4 lb (without magazine). Magazine straight but inserted on the left side

Can be seen at the Museum of Aberdeen Proving Ground, Md

Short recoil-operated, air-cooled MG weighing 34 lb (with feed). Rate of fire 1200-1350 rpm and max vel 2570 ft/sec. Used 7.92 mm German Service ammunition

Was used on the Russian front. Its improved version appeared in 1945 on the Western front under the designation MG-43. It was practically identical with MP-44 described below

Can be seen at the Museum of Aberdeen Proving Ground, Md

Same as above

Gas-operated, air-cooled weapon of remarkable design and mass. It was practically identical with Maschinenpistole 43 (MP-43) and Karabiner 44 (K-44). Overall length 36", barrel ca 19", wt (not given), capacity 30 cartridges of special design

Short, parabolic action rifle, made with the intention of issuing it to civilians for home defense. Overall length 43", barrel 23.2", wt 8.3 lb and magazine capacity 10

Weapon of very original design and of great simplicity. Overall length 34.9", barrel 14.9", wt 9.4 lb and magazine capacity 50

Operated by rapped blow-bolt and cooled by water/air. 400 ft/sec max vel 1875 ft/sec and rate of fire 400-450 rpm

References

11, p 430 and Ref 12

12

10, v 7, pp 187-8 & 11, pp 432 & 438 and Ref 12

4, pp 111-13; 10, v 7, pp 189-91; 11, pp 435-7 & Ref 12

10, v 7, pp 189-197 & 11, pp 439-43

4, pp 176-79; 5, v 1, p 400-91; 11, p 444 and Ref 12

12

11, p 176-91; v 1, pp 444 & 462; 11, pp 509-10 & Ref 12

11, pp 300 and 302

12

11, pp 499-501 and Ref 12

11, pp 300 and 302

12

11, pp 499-501 and Ref 12

11, pp 300 and 302

12

10, v 7, pp 181-3; 11, p 431 and Ref 12

10, v 7, pp 198-9 & 11, pp 445-7

8, v 1, pp 228-31

(SUBMACHINE GUNS AND MACHINE GUNS)



Caliber and Designation

9 mm (.354) Luger (Pamellum)
Automatic Pistol Model 1902
1902/06, 1904 and 1904/06 (M 02,
M 02/06, M 04 and M 04/06)

[See also Note given under 7.65 mm Luger (Pamellum) Pistols M 1900 and 1900/06]

9 mm Luger (Pamellum) Automatic
Pistol Model 1908 (Official German
9 mm Weapon of last WW I; it was
slightly modified in 1920)

Note: Special 6" and 10" barrels were provided for this pistol. The model using an 8" barrel and called 9 mm Pamellum
M 08 Lang (long) was issued to artillery and "Z" host personnel

9 mm Mauser Automatic Pistol,
Military Model, also called
Maschinenpistole. Used in WW I
and to a limited extent in WW II

9 mm Bergmann Automatic Pistol
M 1910, was issued for the Coast
Army. There was also a Model 18-1

9 mm Bergmann Automatic Pistol
(Maschinenpistole) M 1914, called also
Submachine Gun

Note: This weapon was officially adopted by Sweden in 1917 and for this reason is briefly described in the Swedish section.

9 mm Serrv Automatic Pistol,
invented prior to WW I

9 mm Serrv-Falchens Automatic
Pistol (Maschinenpistole) MP,
called in the U.S.A. Submachine
Gun and in Gt Britain Machine
Carbine. Also designated as SI-100

9 mm Walther Automatic Pistol,
invented before WW I

9 mm Walther Automatic Pistol, originally
introduced as Model HR, was officially
designated as P-18. This model was
called "Walther Ammer Pistol"

Note: Several factories made it during WW II

9 mm Schmeisser Machine Pistol,
MP-28 II

9 mm Schmeisser Maschinen Pistole 38
(MP-38), called in the U.S.A. Sub-
machine Gun, Paratrooper Model

9 mm Submachine Gun, MP-34/1,
Bergmann

9 mm Machine Carbine, M-35/1

9 mm Schmeisser Maschinen Pistole
(MP-38) called in the U.S.A. Sub-
machine Gun and Swamp Gun

9 mm Automatic Browning Pistol,
in 1935 designed 10 years earlier
by J.M. Browning. Was used during
WW II by SS troops

9 mm Dreyse Automatic Pistol,
Military Model

9 mm Erbe Machine Pistol, sometimes
called the Schmeisser Machine
Pistol or Carbine

9 mm Johanssens Machine Pistol

9 mm Submachine Gun, EMP-40,
and EMP-44

10.15 mm (.407) Norwegian Rifle

Remarks, Uses and Some Characteristics

Barrel lengths: 6" for M 02 & M 02/06
and 8" for M 04 and M 04/06. The last
two models were issued with a leather
holster attached to a wooden stock. The
M 04 was an official German Navy weapon
used during WW I

Recoil-operated. Length: barrel 6" and
overall 8 1/2"; wt 30 oz; magazine capacity
8 cartridges with round or flat point bullets
weighing 110 and 175 grains. Max vel 1040
to 1500 ft/sec

Some design as 7.65 mm Mauser.
Magazine capacity 10 Luger cartridges.
Could be fired with shoulder stock
holster attached to magazine

Similar in size and design to the
Belgian 9 mm Bergmann-Bayard
except that it was lighter (52 oz)

Modification of Model 18-1

Recoil-operated; magazine capacity
8 rounds

Operated by recoil on the blowback
principle. Overall length 32 1/2"; wt 9 1/2 lb;
magazine capacity 30 Pamellum
cartridges. Max vel 1400 to 1600 ft/sec

Blowback-operated. Served as the prototype
for later models. Capacity 8

Operated by short recoil. Length barrel
4 1/2" and overall 8 1/2"; wt 14 oz; magazine
capacity 8 Pamellum cartridges

Operated by blowback. Overall length (with
stock extended) 35"; wt (without magazine)
9 lb. Magazine capacity 32 Pamellum
cartridges

Can be seen at the Museum of Aberdeen
Proving Ground, Md

Same as above

Slight modification of MP-38; same
dimensions. Cyclic rate of fire 500 rpm

Recoil-operated; length: barrel 6" and
overall 7 1/2"; wt 35 oz; capacity 11

One of the earliest blowback operated
pistols, made in the closing years of WW I

Overall length 33 1/2"; wt 9 lb and cyclic
rate of fire 520 rpm

Capacity 40 cartridges; wt of pistol 9 lb 2 oz

Can be seen at the Museum of Aberdeen
Proving Ground, Md

Used Norwegian ball ammo, type 322

References

4, pp 271-3; 10,
v.1, pp 182 &
417-18 and Ref 12

10, v.1, pp 182 &
417-18; 11, pp 456-
63 and Ref 12

4, pp 275-8; 10,
v.1, p. 420 and
Ref 12

10, v.1, pp 430-41;
11, p. 491 and Ref 12

11, pp 491-2 and
Ref 12

2, p. 322

4, pp 246-8; 11,
pp 456-7 and
Ref 12

2, p. 322

2, p. 322; 4, pp 278-
80; 10, v.1, pp 425-
256; 11, pp 455-55
and Ref 12

11, p. 493 and
Ref 12

11, pp 486-9
and Ref 12

12

12

4, pp 248-50;
7, p. 175; 11, pp 490
and Ref 1

10, v.1, pp 404-
8

10, v.1, pp 408-
10

11, p. 493

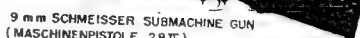
11, p. 494

12

14, p. 8

WEAPONS

(SUBMACHINE GUNS AND MACHINE GUNS)

9 mm BERGMANN SUBMACHINE GUN
(MASCHINENPISTOLE 34/1)9 mm SCHMEISSER SUBMACHINE GUN
(MASCHINENPISTOLE 28 II)9 mm SUBMACHINE GUN
(MASCHINENPISTOLE 38)9 mm SUBMACHINE GUN
(MASCHINENKARABINER 35/1)9 mm SUBMACHINE GUN
(EMP-44)9 mm SUBMACHINE GUN
(MASCHINENPISTOLE EMP-40)15 mm
MACHINE GUN
MG-151/15
(TRIPLE ANTI-AIRCRAFT
GUN ON PEDESTAL
MOUNT)43 mm RHEINMETALL AIRCRAFT
MACHINE GUN MODEL 131, FIXED13 mm RHEINMETALL AIRCRAFT
MACHINE GUN MODEL 131, FLEXIBLE

(Weapons) (cont'd)

Caliber and Designation

- 20 mm Rheinmetall Automatic AA Cannon, Flak 30, developed before WW II
- 20 mm Oerlikon Short Case AC Cannon (2 cm Oerlikon MG-FF)
- 20 mm Oerlikon Automatic AC Cannon, Models F and S, developed by the Oerlikon Co. Zürich and adopted by the Germans before WW II
- 20 mm Oerlikon-AA Cannon (2 cm Flak 29)
- 20 mm Messer Automatic AC Cannon, Model 151 (MG-151), developed before WW II by the Waltherwerk Messer A-G
- 20 mm Messer Automatic AA Cannon, Flak 38
- 20 mm Dutch A/T Rifle [2 cm PaB 785 (S)]
- 20 mm Fanch Machine Gun [2 cm MG 39 (L)]
- 20 mm Solothurn Cannons:
2 cm KwK 30, 2 cm KwK 38
2 cm Flak 30, 2 cm Flak 38
2 cm Flak Victoria 38
2 cm GdFlak 38 and Italian
2 cm M 35 (L)
- 20 mm Messer Machine Gun, MG-115, developed during WW II
- 20 mm Recoiless Cannon (9 barrels)
- 20 mm and 25 mm Semag Automatic Cannon for Infantry (Mounted on a wheeled carriage)
- 25 mm (.98P) French AA Gun [2.5 cm Flak Hotchkiss (D)]
- 25 mm French A/T Guns: 2.5 cm PaB 122 H 113 (D) and 2.5 cm KwK 121 (D)
- 27 mm (1.06P) Signal Pistol (Kampfpistol), Modified
- 78/28 mm (1.102/0.78P) Tapered Bore A/T Rifle (AP-Rifle 41), called also Squeeze Bore or Gerlich Gun
- 30 mm (1.181P) Messer Machine Gun, MK-213C, developed during WW II
- 30 mm Rheinmetall Automatic AC Cannon, MK-101, developed in 1942
- 30 mm Rheinmetall Automatic AC Cannon MK-103, developed in 1943
- 30 mm Rheinmetall Automatic AC Cannon MK-108, developed in 1944
- 30 mm Automatic Recoiless Cannons, SC-16, SC-17 and SC-18, developed during WW II by the H.Göting Werke
- 30 mm Solothurn AC Cannon (3 cm Flak K)
- 30 mm Aircraft Machine Cannon, MK-303

Remarks, Uses and Some Characteristics

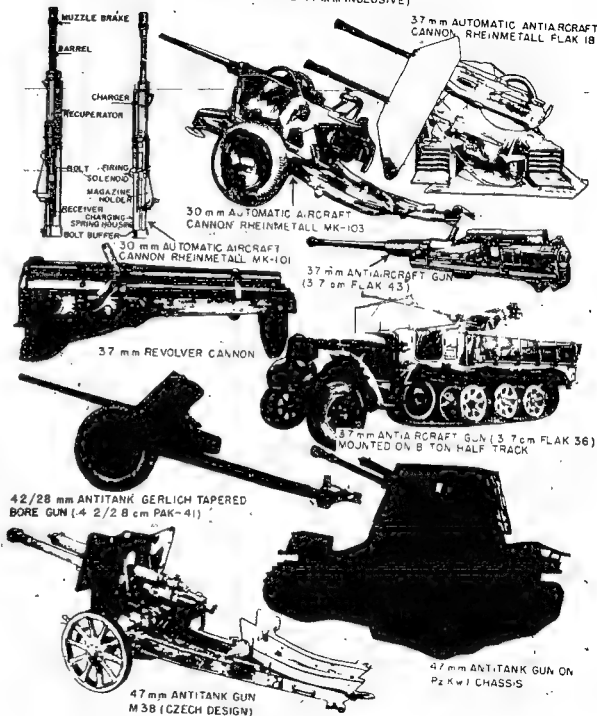
- Short recoil-operated and air-cooled. Wt (with feed) 141 lb, rate of fire 200-280 and max vel 2950 ft/sec. Used HE-T projectiles.
- Used projectiles: HE, HE (self-destruying), HE-T, AP, APHE and AP.
- Blowback-operated and air-cooled. Wt (with feed) 150 lb, rate of fire 180 and max vel 2610 ft/sec
- Used AP, AP-T, HE, HE-T, HE-T-T and HE-T self-destruying projectiles
- Can be seen at the Museum of Aberdeen Proving Ground, Md
- Short recoil-operated and air-cooled. Wt (with feed) 92½ lb, rate 700-750 and max vel 2900 ft/sec. Called by South (Ref 9) one of the most remarkable AC MGs in existence
- Short recoil-operated and air-cooled. Wt (with feed) 129 lb, rate of fire 420-480 and max vel 2950 ft/sec
- Used Dutch AP and HE ammo
- Used French HE shell, type 39
- Used ammunition:
HE, HE-T, HE-T-T
HE-T, HE-T (self-destruying), HE (Italian), AP, AP-T, AP-T-T, AP-T (self-destruying), AP-T (Italian) and AP (Italian)
- Not described here because the reference is confidential
- Can be seen at the Museum of Aberdeen Proving Ground, Md
- Developed in 1921 and 1923 but not adopted in Germany because it was considered to be too heavy. A number of Semags were sold before 1930 to China and to Spain
- Used French HE and HE-T shells
- Used French AP type 114 shell
- Can be seen at the Museum of Aberdeen Proving Ground, Md
- Used ammo: HE (2.8 cm Sprgr Patz 41) and AP (Patz Patz 41)
- Not described here because Ref 8, v 3 is confidential
- Short recoil-operated and air-cooled. Wt (with feed) 335 lb, rate of fire 230-260 and max vel 2950 ft/sec
- Operated by gas-actuated piston and air-cooled. Wt (with feed) 308 lb, rate of fire 420 and max vel 2820 ft/sec
- Blowback-operated and gas-cooled. Wt (with feed) 135 lb, rate of fire 400-450 and max vel 1640 ft/sec
- Not described here because the reference is confidential
- Used HE and AP ammo: 3 cm Sprgr and 3 cm Patz 40
- Can be seen at the Museum of Aberdeen Proving Ground, Md

References

- 5a, table 1 and 8, v 1, p 666
- 5a, pp 44-5
- 5a, p 44; 8, v 1, pp 516 & 618 and Ref 12
- 5a, p 43
- 12 =
- 5a, pp 45; 8, v 1, pp 602-4 & 606; 11, p 501 and Ref 12
- 8, v 1, pp 605-6 & 666 and Ref 12
- 5a, p 13
- 5a, p 13
- 5a, pp 43-4
- 8, v 3, pp 46-51
- 12
- 8, v 1, pp 514-15
- 5a, p 14 and Ref 12
- 5a, p 14
- 12 =
- 5a, p 14; 9, p 371 and Ref 12
- 8, v 3, p 44
- 8, v 1, pp 555-61 & 666-8
- 8, v 1, pp 555-61 & 666-8 & Ref 12
- 5a, p 44
- 8, v 4, pp 639-31
- 9, p 379
- 12

WEAPONS

(CALIBERS 30 mm to 4.7 mm INCLUSIVE)



(Weapons) (cont'd)

Caliber and Designation

Remarks, Uses and Some Characteristics

References

37 mm (1.457") Rheinmetall Automatic A/T Cannon, Type 18 (3 cm Flak 18), developed prior to WW II by Rheinmetall-Borsig A-G

Short recoil operated and air-cooled, Wt (with feed) 555 lb, rate of fire 140-180 and max vel 2120 ft/sec. Used projectiles: HE, HEI, HEI-T, HE (high capacity) and AP

8, v.1, pp 554 & 666;
7a, pp 45-6 & 9, p 384

37 mm A/T Cannon, 3.7 cm Flak 36, Flak 37 and Flak 41

Used ammo HE (3.7 cm SprgPatr 18), HE, high capacity (SprgPatr 18), HEI (B/SprgPatr 18), HEI-T (B/SprgPatr 18 L'spur) and AP, without cap (SprgPatr 18)

7a, pp 45-6 and 9, p 384

37 mm A/T Cannon (3.7 cm Pak)

Used: AP proj with core, arrowhead design (3.7 cm SprgPatr 40); AP proj without cap (SprgPatr 37) and HE proj 18 modified (SprgPatr 18 mod)

5a, p 15 and 9, pp 373 & 386

37 mm A/T Cannon, Fixed Defense (3.7 cm Pak K)

Used ammo: HE (3.7 cm SprgPatr) and AP (SprgPatr 18 mod)

5a, p 15

37 mm Naval Gun: 3.7 cm SK C/30

Used ammo: HE (3.7 cm SprgPatr 40) and HE-T (SprgPatr 18 L'spur)

5a, p 15 and 9, pp 382 & 388

37 mm Naval Gun: 3.7 cm SK C/36

Used HE projectiles

5b, table 1

37 mm Tank Gun: 3.7 cm KwK

Used ammo: HE (3.7 cm SprgPatr 18 mod & SprgPatr 40), HE-T (SprgPatr 18 L'spur), AP (SprgPatr 18 mod & SprgPatr 40) and Stick grenade (Steiger 41)

5a, p 35

37 mm A/T Gun: 3.7 cm Pak 41

Used stick (modified) bomb, 3.7 cm Steiger 41

9, p 383

37 mm Czech A/T Gun: 3.7 cm

Used Czech ammo: HE (3.7 cm SprgPatr 34), AP (SprgPatr 34, 37, 37 mod & 40/37) and Stick Grenade (Steiger 41)

9, p 316

37 mm Czech Tank Gun: 3.7 cm KwK 38 (t)

Same as above

5a, p 36

37 mm Czech Tank Gun: 3.7 cm KwK 38 (t)

Used French HE and AP ammo: 3.7 cm SprgPatr 143, 147, 148 (t) and PzSprgPatr 145 & 146 (t)

5a, p 35

37 mm Czech Tank Gun: 3.7 cm KwK 38 (t)

No description given

5a, p 39

37 mm Czech Tank Gun: 3.7 cm KwK 38 (t)

No description given

5a, p 39

37 mm Russian Infantry Howitzers: 3.7 cm IG 143 & 146 (t)

No description given

9, p 382

37 mm Polish A/T Gun, called by the Germans 3.7 cm Pak (p)

Used Polish design AP proj: 3.7 cm Pzgr (p)

12

37 mm Cannon: Flak 36, Pak 37, Flak 43, Kavalier Cannon and AC Cannon (used on Stuka aircraft)

Can be seen at the Museum of Aberdeen Proving Ground, Md

12

40 mm (1.575") AA Gun, Type 18 (4 cm Flak 18)

Used ammo: HE (4 cm SprgPatr 18), HE-T (SprgPatr 18 L'spur), HEI (B/SprgPatr 18 L'spur) and AP-T (PzSprgPatr 18 L'spur)

5a, p 46 and 9, pp 388-9

42 mm (1.654") Tapered Bore Gun 41 (4.2 cm Pak 41), called also Griffling Gun or Squeeze Bore Gun

Used ammo: HE (4.2 cm SprgPatr 41) and AP with core (PzSprgPatr)

5a, p 46 and 9, pp 388-9

43.3 (1.75") mm Recoilless Grenade Discharger Panzerfaust 30, also known as (Armored Fist, type 30, small) formerly called Panzerfaust 2 (First Cartridge, type 1) and a larger model Panzerfaust 30, formerly called Panzerfaust 2

Smooth-bore tube, 1.75" diameter and 31.5" long which fired a hollow charge/TNT missile, resembling to appearance a rounded hand grenade. Projectile available at Museum of Picatinny Arsenal (100') long of which the warhead is 9.5" long and the lined cylindrical body is 10". Diameter of warhead 1.75" and of body 1.75"

9, pp 339-40
11, p 522
and Ref 13

45 mm (1.77") Russian A/T Guns 45 cm Pak 124 & 124.1 (t)

Used Russian HE and AP ammo

5a, p 17

45 mm Russian Tank Guns, 4.5 cm KwK 184 2, 184/3 & 184 4 (t)

No description, given

5a, p 59

45 mm Russian Infantry Howitzers: 4.5 cm IG 186 (t)

Used HE bomb, Wgr (t)

5a, p 26

45 mm Italian Mortar: 4.5 cm GrW 176 (t)

No description given

5a, p 26

46 mm (1.811") Polish Mortar: 4.6 cm GrW 13 (p) and GrW 16 (p)

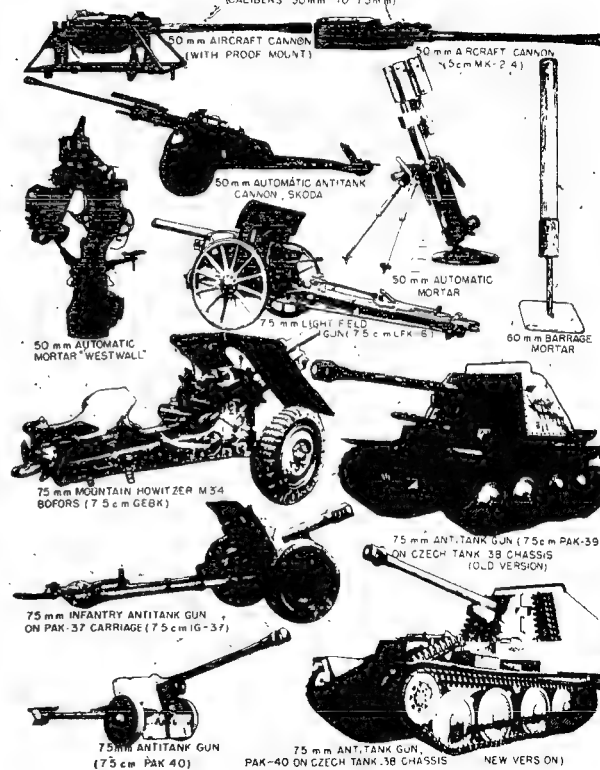
No description given

5a, p 26

ver 242

WEAPONS

CALIBERS 50mm TO 75mm



Caliber and Designation

47 mm (1.85") Austrian "Böhler" Gun
(4.7 cm Böhler (G)) or 4.7 cm Pak
Böhler (G)

47 mm Belgian A/T Gun (4.7 cm
Pak 185 (B))

47 mm Czech Guns: 4.7 cm K 36 (t),
Pak Skoda 1936 (t) and Flak 37 (t)

47 mm French A/T Gun (4.7 cm
Pak 180 u 183 (D))

47 mm French Tank Gun: 4.7 cm
KwK 173 (B)

47 mm Italian A/T Gun (4.7 cm
Pak 177 (I))

50 mm (1.9685") Tank Gun:
5 cm KwK

50 mm Tank Gun 38: 5 cm KwK 38

50 mm Long Tank Gun: 5 cm KwK 39
(L/60), KwK 39:1 and KwK 39/2 (L/60)

50 mm Tank Gun: 5 cm KwK 40
and KwK 42

50 mm A/T Gun 38 (5 cm Pak 38)

Note: According to Ref 5b table 1, this gun existed in 50 and 60 caliber lengths and was designated as 5 cm Pak (L/50)

50 mm A/T Casemate and Turret
Gun, long mount, 5 cm Pak K&T
(K&L)

50 mm A/T Casemate and Turret
Gun, short mount, 5 cm Pak KvT
(K&L)

50 mm Light Mortar: 5 cm GrW 36
and GrW M 19

50 mm AA Gun 41 (5 cm Flak 41)

50 mm Automatic Aircraft Cannon
(5 cm BK) developed during WW II
by the Rheinmetall-Borsig A-G

50 mm Automatic AC Cannon, MK-214

50 mm AC Cannon

50 mm A/T Automatic Cannon, Skoda

50 mm Automatic Mortar (Westwall)

50 mm Belgian Light Mortar: 5 cm
GrW 203 (t)

50 mm French Light Mortar: 5 cm
GrW 203 (t)

50 mm Russian Light Mortar: 5 cm
GrW 203 (t)

50.8 mm (2") British Mortar: 5 cm
GrW 202 (e)

55 mm (2.165") Aircraft Automatic
Cannon, MK-112, developed near
the end of WW II by the Rhein-
metall-Borsig A-G

55 mm Automatic Cannon, MK-114,
not fully developed during WW II

55 mm Automatic Recoilless
Cannon, MK-115, developed by
Rheinmetall-Borsig A-G but
not put into production

(Weapons) (cont'd)

Remarks: Uses and Some Characteristics
Used Austrian design AP and HE ammo;
4.7 cm Sprg Patr 35 (G) and Sprg Patr (G)
Böhler (G)

Used Belgian HE and AP ammo

Used Czech design HE and AP ammo;
4.7 cm Sprg Patr 36 (t), Sprg Patr 36 (t)
and Flak 37 (t)

Used French HE and AP ammo: 4.7 cm
Sprg Patr and Sprg Patr

Used French HE and AP ammo: 4.7 cm
Sprg Patr 173 (G) and Sprg Patr 176 (t)

Used Italian HE and AP ammo

Used ammo: HE (5 cm Sprg Patr 38), AP
(Sprg Patr 39, 40 & 40/1) and Stick Grenade
(Steigler 42)

Used AP ammo: 5 cm Sprg Patr

Used ammo: HE (5 cm Sprg Patr 38), AP
(Sprg Patr 39, 40 & 40/1) and Stick Grenade
(Steigler 42)

Used ammo: HE (5 cm Sprg Patr 38), AP
(Sprg Patr 39, 40 & 40/1) and Stick Grenade
(Steigler 42)

Used ammo: HE (5 cm Sprg Patr 38) and
stick grenade (Steigler 42)

Used ammo: HE (5 cm Sprg Patr 38), AP
(Sprg Patr 39, 40 & 40/1) and stick grenade
(Steigler 42)

Used ammo: Short HE (Ka 5 cm Sprg Patr 38)
and Short AP (Ka 5 cm Sprg Patr (Pak KvT (K&L))

Used HE mortar ammo such as: 5 cm Vpr Patr
36, 39 & 41

Used ammo: HE-T (5 cm Br Sprg Patr 41
L'apuri), HE-T (Sprg Patr L'apuri), AP (Sprg Patr
19 & 43) and AP (Sprg Patr 42 V)

No description is given here because Ref 8, v3
is confidential

Can be seen at the Museum of Aberdeen
Proving Ground, Md

Same as above

Same as above

Same as above

Used various mortar ammo: Belgian, French,
German and Russian

Same as above

Same as above

Same as above

Used British HE and smoke bombs

Not described here because Ref 8, v3 is
considered confidential

Same as above

Same as above

Same as above

Same as above

References

8, v3, p 17 and
9, pp 391-2

8, v3, p 17

8, v3, p 18, 9,
pp 390-2 & Ref 12

8, v3, p 17

8, v3, p 36

8, v3, p 17

8, v3, pp 36-7 and
9, pp 376 & 395-5

9, pp 395 & Ref 12

8, v3, p 37

8, v3, pp 36-7

8, v3, p 18

8, v3, p 19

8, v3, p 19

8, v3, pp 6-7 and
9, pp 390-1

8, v3, p 46 and
9, p 395

8, v3, p 638

12

12

12

12

12

12

12

8, v3, p 26

8, v3, p 26

8, v3, p 26

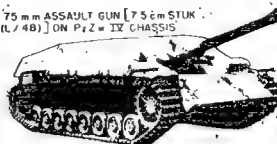
8, v3, p 27

8, v3, pp 614 &
627

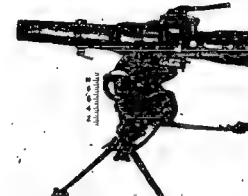
8, v3, p 636

8, v3, p 637

75 mm ASSAULT GUN (7.5 cm STUK
L/48) ON Pz 2 w IX CHASSIS



75 mm GERLICH TAPERED BORE
ANTI-TANK GUN (7.5/55 cm PAK-41)



75 mm RECOILLESS GUN
(7.5 cm LG-40) ON TRIPOD



75 mm PANTRY HOWITZER
(7.5 cm HK-42) SMOOTH BORE



75 mm ANTITANK GUN PAK-42
(7.5 cm PAK 40 ON LFH 18/40 CARRIAGE)



75 mm RECOILLESS GUN
(7.5 cm RPK-43)



75 mm ASSAULT GUN (7.5 cm STUK-42)
ON Pz 39g IV CHASSIS

75 mm ANTITANK GUN (7.5 cm PAK 50)
ON PAK 38 CARRIAGE (EXPERIMENTAL)

(Weapons) (cont'd)

Caliber and Designation	Remarks, Uses and Some Characteristics	Reference
60 mm (2.362") French Mortar 6 cm GrW 225 (f)	Used French HE cast steel bomb: 6 cm Stg (Schikgus) Wgr 225 (f)	5a, p 27
60 mm Mortar Damage	Can be seen at the Museum of Aberdeen Proving Ground, Md	12
63 mm (2.559") French Mountain Pack Howitzer: 6.5 cm GebK 221 (f)	Used French HE shell: 6.5 cm Gr/Pz (f)	5a, p 52
65 mm French Quick-Firing Gun: 6.5 cm Schenk Schnellartillerie (f) (2)	Used French ammo: HE [6.5 cm Gr/Pz A2, 6.5 cm Schenk Schnellartillerie (f)] and AP [Gr/Pz (f)]	5a, p 60
65 mm Italian Mountain Pack (Howitzer): 6.5 cm GebK 716 (f)	Used Italian ammo: HE [6.5 cm Sprg/Pz (f)] and AP [Gr/Pz (f)]	5a, p 52
65 mm Yugoslav Mountain (Pack) Howitzer: 6.5 cm GebK 222 (f)	Used Yugoslav ammo: HE [6.5 cm Sprg/Pz 221 (f)] and AP [Sch/Pz 221 (f)]	5a, p 52
73 mm (2.874") Rocket Launcher, Falm Gwrt, capable of firing 35 rockets simultaneously	A 15-frame launcher with fast elevating and traverse gears. It fired 7.5 cm Raketenpropagranate or 7.5 cm Propagranatepropagranate 41	9, pp 23-46
75 mm Mountain Gun: 7.5 cm GebK 15, 16 & GebK 14/15	Used ammo: HE (7.5 cm GebK 15, GebK 15 A1, GebK 15 R, GebK 39), HoC (Gr 39 H/A), as well as some Austrian and Czech ammo	5a, pp 35 and 9, pp 399-403
75 mm Glade Mountain Gun M 15: 7.5 cm GebK M 15	Same ammo as above	5a, p 35
75 mm Light Field Gun 16: 7.5 cm IFK 16	Can be seen at the Museum of Aberdeen Proving Ground, Md	12
75 mm Field Gun 16/1: 7.5 cm FK 16/1	Used HE proj (7.5 cm KGrGrKPS) and AP proj (KGrGrKPS)	9, pp 421 & 423
7.5 cm Field Gun 16A, new pattern: 7.5 cm FK 16A	Used same ammo as above, plus HoC proj (7.5 cm G15 H/A)	5a, pp 60-1 and 9, pp 409, 421 & 423
75 mm Light Field Gun 18: 7.5 cm IFK 18	Used ammo: HE (7.5 cm Sprg/Pz 34 & KGrGrKPS), AP (KGrGrKPS), APC (Gr/Pz 34), HoC (Gr 38 H/A & Gr/Pz 38 H/A) and Smoke (Nbr/Pz)	5a, pp 41-2 and 9, pp 400-3, 407, 409, 421 & 423
75 mm Light Infantry Howitzer 18: 7.5 cm IFK 18	Can be seen at the Museum of Aberdeen Proving Ground, Md	12
75 mm Light Mountain Infantry Howitzer 18: 7.5 cm GebK M 18	Used ammo: HE (7.5 cm Jgr 38 FES) and HoC (Jgr 38 H/A and H/B)	9, pp 413, 418 & 419 and Ref 12
75 mm Light Infantry Gun: 7.5 cm IFK 18, 37 & 42	Used same projectiles as previous weapon	9, pp 413, 418 & 425
75 mm Light Infantry Mountain Gun: 7.5 cm GebK G 18	Used ammo: HE (7.5 cm Jgr 18 A4, HoC (Jgr 38 H/A, Jgr 38 H/B), Jgr/Pz H/A, Jgr 38 H/B) and incendiary shell (Jgr Dru)	5a, p 30; 9, pp 404 and Ref 12
75 mm Heavy Heavy Gun: 7.5 cm s/G 35, s/G 35/1 & s/G 43	Same as above	5a, p 30 and 9, pp 404-5
75 mm Naval Gun: 7.5 cm SK C/F (L/33)	Used ammo: HE (15 cm Jgr 33, Jgr 38 & Jgr 19 A4), HoC (Jgr 39 H/A & Jgr 39 H/B), Smoke granule (Gwrt 42), Smoke (Jgr 38 H/B) and Inc (Jgr 38 B)	5a, p 31
75 mm Mountain Howitzer Boomer: 7.5 cm GebK 54	Used HE projectiles	5a, table 1
75 mm Mountain Gun 36: 7.5 cm GebK 36	Can be seen at the Museum of Aberdeen Proving Ground, Md	12
75 mm Gun 37: 7.5 cm K 37 L/24	Used ammo: HE (7.5 cm Gr 34 Sprg/Pz 34, KGrGrKPS & KGr 34 A4), HoC (Gr 38 H/A & H/B) and Smoke (Nbr/Pz) and some foreign ammo	5a, pp 32 and 9, pp 396, 401, 409 & 416
75 mm Tank Gun: 7.5 cm KwK	Used ammo: HE (7.5 cm Sprg/Pz), HoC (Gr/Pz 38 H/A, H/B & H/C), AP (Gr/Pz), Case Shot (Gr/Pz), Smoke (Nbr/Pz) and incendiary shell (KGrGrK Ror Dru)	5a, p 58
75 mm Assault Gun: 7.5 cm StuK	Same as above	5a, p 38
75 mm Field Gun 38: 7.5 cm FK 38	Used ammo: HE (7.5 cm KGrGrK, Sprg L/4.0), HoC (Gr/Pz 38 H/B & H/C) and Smoke (Nbr/Pz) (KGrGrK Ror Dru)	5a, p 62 and 9, p 415

(Weapons) (Cont'd)

Caliber and Designation	Remarks, Uses and Some Characteristics	Reference
75 mm Tank Gun 38: 7.5 cm KwK 38	Used HoC ammo: 7.5 cm Gr/Pz 38 H/A	9, p 409
75 mm A/T Gun 39: 7.5 cm Pak 39 L/48	Used ammo: HE (7.5 cm Sprg/Pz 34), HoC (Gr/Pz 38 H/A, H/B & H/C), AP (Gr/Pz 39, Gr/Pz 40 & Gr/Pz 40 W) and Smoke (Nbr/Pz)	5a, p 39
75 mm Tank Gun: 7.5 cm KwK 40 L/43 and KwK 40 L/48	Same as above	5a, p 39
75 mm Assault Gun: 7.5 cm StuK L/43 & StuK L/48	Same as above	5a, p 39
75 mm A/T Gun: 7.5 cm Pak 97/38 and 97/40	Used ammo: HE (7.5 cm Sprg/Pz), HoC (Gr/Pz 15/38 H/A, Gr/Pz 38 H/A, Gr/Pz 38/97 H/A & H/B), AP (Gr/Pz 39), and StuK (Gr/Pz 39, Gr/Pz 40 & Gr/Pz 40 W) and Smoke (Nbr/Pz)	5a, p 21 and 9, pp 215, 419-20 & 425
75/50 mm Skoda Dual Purpose Gun	Used HE ammo: 7.5 cm Sprg/Pz 34 & Sprg/Pz 34 APC (Gr/Pz 34), HoC (Gr/Pz 38 H/A, H/B & H/C), AP (Gr/Pz 40, Weichschuss or Gr/Pz 40, barter Kern) and Smoke (Nbr/Pz)	5a, p 21; 9, p 406
75 mm A/T Gun 40: 7.5 cm Pak 40	Used ammo: HE (7.5 cm Sprg/Pz 34 KwK, etc), HoC (Gr/Pz 38 H/A, H/B & H/C), AP (Gr/Pz 40, Weichschuss or Gr/Pz 40, barter Kern) and Smoke (Nbr/Pz)	5a, p 21; 9, pp 398, 401-2, 409-9, 411 & 417 & Ref 12
75 mm Self-Propelled A/T Gun: 7.5 cm Pak 40/1 (S), Pak 40/2 (S) and Pak 40/3 (S)	Used HoC ammo, such as 7.5 cm Gr/Pz H/B	5a, p 21; 9, p 411 and Ref 12
75 mm Tank Gun 40: 7.5 cm KwK 40	Used ammo: HE (7.5 cm Sprg/Pz 34 & Sprg/Pz 34 APC (Gr/Pz 34), HoC (Gr/Pz 38 H/A, H/B & H/C), AP (Gr/Pz 40, Weichschuss or Gr/Pz 40, barter Kern) and Smoke (Nbr/Pz)	9, pp 398, 400-3, 409, 411 & 417
75 mm Recoilless Gun for Airborne Troops, Type 40 (7.5 cm Leichter Geschütz 40)	Used same ammo as above, plus Sprg 34 and Gr 38 H/B	9, pp 398, 400-3, 409 H/A Ref 12
75 mm Assault Gun 40 (7.5 cm StuG 40)	Used ammo: HE (7.5 cm Sprg/Pz 34), APC (Gr/Pz 34), HoC (Gr/Pz 38 H/A & H/B), AP (Gr/Pz 40, Weichschuss or Gr/Pz 40, barter Kern) and Smoke (Nbr/Pz)	9, pp 398, 400-2 & 409 H/A
75 mm Assault Gun: 7.5 cm StuK 40 L/43 and StuK 40 L/48	Used ammo: HE (7.5 cm Sprg/Pz 34) and HoC (Gr 38 H/B)	9, pp 413 & 417 and Ref 12
75/55 mm A/T Gun 41: 7.5/5.5 cm Pak 41 (Gr/Pz Type Gun, called also Tapered Bore Gun, Reducing Bore Gun or Squeeze Bore Gun)	Can be seen at the Museum of Aberdeen Proving Ground, Md. Used AP proj with iron core (7.5 cm Pak 41 W) and AP proj with tungsten carbide core, arrowhead design (Gr/Pz 41 H/K)	5a, p 21 and 9, pp 378 & 408 and Ref 12
75 mm Assault Gun 42: 7.5 cm StuK 42	Used ammo: HE (7.5 cm Sprg/Pz 42), HoC (Gr/Pz 38 H/B & AP (Gr/Pz 39/42, 40 & 40/42)	5a, p 39
75 mm Tank Gun 42: 7.5 cm KwK 42 L/70	Same as above	5a, p 39
75 mm Tank Gun 42: 7.5 cm KwK 42	Used ammo: HE (7.5 cm Sprg/Pz 42) and AP (Gr/Pz 39/42)	9, pp 411 & 423 and Ref 12
75 mm Assault Gun 42: 7.5 cm StuK 42 L/70	Used same ammo as above	9, pp 411 & 423 and Ref 12
75 mm Infantry Howitzer 42, Smooth Bore: 7.5 cm IFK 42	Can be seen at the Museum of Aberdeen Proving Ground, Md	12
75 mm Recoilless Gun 43: 7.5 cm RFG (Rückstoßfreie Kanone) 43	Can be seen at the Museum of Aberdeen Proving Ground, Md. Used HoC proj: 7.5 cm Gr/Pz 43 H	5a, p 21 and Ref 12
75 mm A/T Gun 50, Experimental: 7.5 cm Pak 50	Can be seen at the Museum of Aberdeen Proving Ground, Md	12
75 mm Belgian Gun: 7.5 cm FK 234 (b)	Used ammo: HE: Sprg 230/7 (f) and HoC: Gr 15/38 H/B (f)	5a, p 21 and 9, pp 415, 420-1 & 425
7.5 cm FK 233 (b)	HE: Sprg 240/7 (b)	
7.5 cm FK 236 (b)	HE: Sprg 1900/15 (b)	
75 mm Czech AA Gun: 7.5 cm Flak (Stoak)	Used Czech HE ammo: 7.5 cm Sprg/Pz (a)	5a, p 46
75 mm Czech Field Gun 17: 7.5 cm FK 17 (f)	Used Czech HE ammo: 7.5 cm Gr M/17 & M/19 (f)	5a, p 66
75 mm Dutch Gun: 7.5 cm FK 243 (b)	Used ammo: HoC: Gr 38 H/C (b)	9, pp 413, 421 & 423
7.5 cm FK 243 (b) L 30	HE: KGrGrKPS and KGr Ror Pa	

(Weapons) (cont'd)

Caliber and Designation

75 mm French Gun:
7.5 cm FK 231 (f), Mle 97
7.5 cm FK 232 (f), Mle 97/13
7.5 cm GekK 234 (f), Mle 1928
7.5 cm GekK 231 (f), Mle 1935
7.5 cm Flak M 17/34 & Flak M 36
75 mm Field Gun: 7.5 cm FK 237 (f)
744 (f)
75 mm Italian Mountain Gun: 7.5 cm
GebK 259 (f)
(See also under Weapons in the Italian section)
75 mm Norwegian Gun:
7.5 cm FK Schneider (a)
7.5 cm FK 01 (a)
7.5 cm FK L/17 (a)
7.5 cm FK 240 & 247 (a)
75 mm Polish Gun:
7.5 cm FK 97 (p)
7.5 cm FK 02/20 (p)
75 mm Yugoslav Gun:
7.5 cm FK 249 (f), Mod 12 (Schneider)
7.5 cm GekK 239 (f)
7.5 cm GekK 239 (f)
7.5 cm GekK 285 (f)
75 mm Yugoslav Mortar:
5 cm GekK 229 (f)
76 mm (2,892) British AA Gun:
7.6 cm Flak (a)
76.2 mm (3,000) Russian Gun:
7.62 cm FK 19 (f)
7.62 cm FK 200/1 and 310 (f)
7.62 cm FK 36 (f)
7.62 cm FK 200 (f)
and many other models were captured
and used by the Germans during WW II
(See Weapons in the Russian section)
76.5 mm (3,000) Austrian Field Gun:
7.65 cm FK 8/18: FK 17b and FK 18b,
manufactured by Skoda Works, Pilsen
76.5 mm French Field Gun:
7.65 cm FK 3/8 (f) & FK 17
76.5 mm Yugoslav Gun: 7.65 cm FK 100
(f), 303 (f), & 304 (f), manufactured by
Skoda Works
77.43 mm (3,031.77) Recoilless
Antitank Cannon, SdKfz 131a,
developed during WW II by the
H.G.ing Werke but not put into
production
80 mm (3,150) Medium Mortar,
designated B cm SdGr 34
80 mm Medium Mortar, designated as
7.5 cm GekK 34
80 mm Trench Mortar, designated as
7.5 cm GekK 42
80 mm Automatic Mortar, "Pilsen"
80 mm A/T Gun (High-Low Pressure
Mortar Bomb Projector)
80 mm Smooth-Bore Weapon, called
Panzerfaustflamme, developed by the
Rheinmetall-Borsig Co. and issued to
the troops at the end of 1944

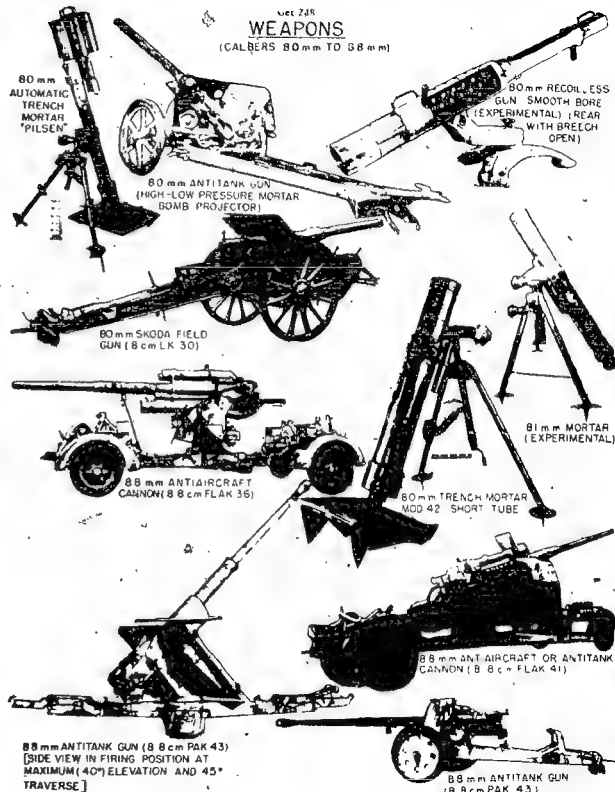
Remarks, Uses and Some Characteristics

Used ammo
HE: Sprgr 1900/15 (f) & Sprgr 231/1 (f) and
HoC: Gr 15/34 H/B (f)
HE: Sprgr 232/1 (f) & Sprgr 264 (f) and
HoC: Gr 15/34 H/B (f)
HE: Sprgr 231 (f)
HE: Sprgr 248 (f)
Used Italian HE and Shrapnel ammo
Lead same ammo as 7.5 cm GebK 25
Lead Norwegian ammo
HE: GekK 231 (a) and Shrapnel: GekK 234
HE: GekK 231 (a) & M/21 & M/36(a) and
HE: GekK 231 (a) & M/21 & M/36 (a); HE: GekK
234 (a)
No information available
Used ammo
HE: Sprgr 1900/15 (f) & HoC: Jgr 38 H/B
Used ammo
HE: Sprgr 264 (f) & Sprgr 1900/15 (f) and
HoC: Gr 15/34 H/B (f) & Gr 34/97 H/C (f)
Same ammo as 7.5 cm GebK 15
HE: Sprgr 249 (f) and Shrapnel (Sch 250 & 25)
HE: Sprgr 260/1 & 260/2 (f)
Used HE bombs: Wgr 229 (f)
Used British HE Bomb round: 7.6 cm
Sprgr Pat (a)
Used various Russian design projectiles
either captured or manufactured in Germany
Used Austrian and Czech design ammo
Used French design ammo
Used Yugoslav, Czech and Austrian ammo
Not described here because Ref B,p 3 is
confidential
Used HE mortar ammo: 8 cm Wgr 34,
Wgr 38, Wgr 39 & Wgr 38 Deva
Lead smoke mortar ammo (7.5 cm Wgr 34 Nb)
Used HE mortar ammo (7.5 cm Wgr 34)
and Smoke Wgr 34NO
Can be seen at the Museum of Aberdeen
Proving Ground, Md
Same as above

References

5a, pp 21 & 41
9, pp 413-25
5a, p 64
5a, p 55
5a, pp 55 &
65-66
5a, p 21 and
9, pp 413-20
& 423
5a, pp 21, 54-5 &
9, pp 413, 419-20
and 423
5a, p 27
5a, p 48
5a, pp 25-4, &
40-1; 9, pp 426-32
5a, p 68
5a, pp 68-9
5a, pp 68-9
8, v 3, p 650
9, pp 529, 531 &
535
9, p 532
9, pp 532-3
and Ref 12
12
12
6, p 188

Let 248
WEAPONS
(CALIBERS 80 mm TO 88 mm)



(Weapons) (cont'd)

Caliber and Designation

40 mm Multiple-Rocket Launcher, designated as 8 cm Raketenw.-faehrer.
80 mm Trench Mortar, Short Tube, Mod 42
80 mm Recoilless Gun, Smooth Bore, Experimental
80 mm Czech Field Gun: 8 cm K 16/17 (9)
80 mm Czech Field Gun: 8 cm FK 30 (1)
80 mm Polish Mountain Gun: 8 cm L/28 (p)
81 mm (3.19") Mortar, Experimental
82 mm Foreign Mortars used by the Germans included: 8.1 cm GrW 274 (dual), 8.1 cm GrW 270 (3), 8.1 cm GrW 280 (3), 8.1 cm GrW 278 (f), 8.1 cm GrW 280 (f), 8.1 cm GrW 274 (f) & 274 (f)
81 mm (3.19") Czech Design AA Gun: 8.35 cm FlaK M/32 (t)
83.9 mm (3.3") British Field Gun: 8.35 cm F.C. 27, 27.2 cm F.C. 27 (a)
83.9 mm Russian Field Gun: 8.35 cm F.C. 305 (t)
86 mm (3.4") Single Barrel Rocket Launcher, designated as 8.6 cm R.G. 42 and weighing 40 kg
86 mm Rocket Launcher (No German designation is given)
87 mm (3.4") British Field Gun: 280, 281 & 282 (a) (25 pounders)
88 mm (3.45") AA Gun 18: 8.8 cm Flak 18
88 mm Tank Gun 36: 8.8 cm KwK 36
88 mm Naval Gun: 8.8 cm SK C/35, C/30/31, C/32 & C/33
88 mm Torpedobomb Gun: 8.8 cm Tbc. R L/45
88 mm Tank Gun 36/36 calibers (dual): 8.8 cm KwK 36 L/45
88 mm AA Gun 30: 8.8 cm Flak 36
88 mm AA Gun 37: 8.8 cm Flak 37
88 mm AA Gun 41: 8.8 cm Flak 41
88 mm AA Gun 41: 8.8 cm Flak 41
88 mm Short Mortar
89 mm Trk. Gun 43: 8.9 cm KwK 43 (L/71)

Remarks, Uses and Some Characteristics

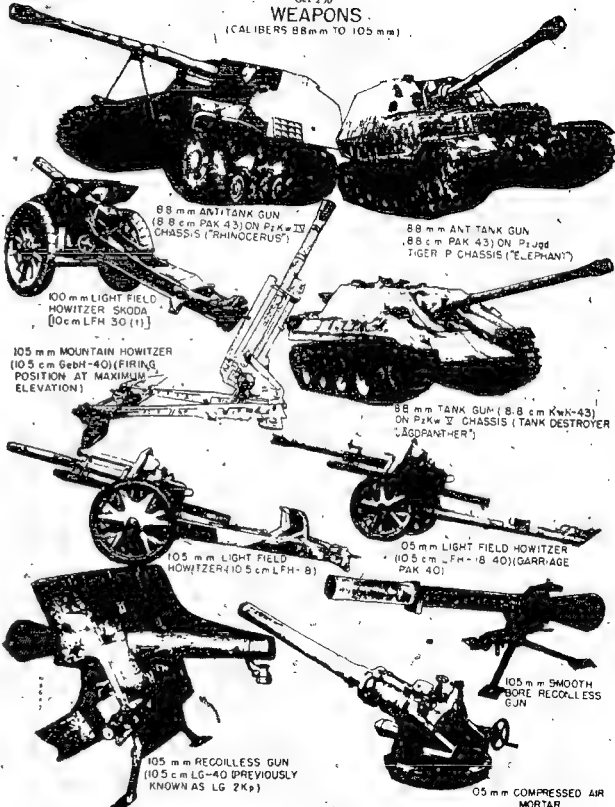
It fired HE aircraft rockets (8 cm Raketen Sprenggranaten), similar in construction to a standard Russian aircraft rocket
Can be seen at the Museum of Aberdeen Proving Ground, Md
Same as above
Used Czech HE shell, 8 cm GrW 30/17 (t)
Used Czech HE and AP projectiles: Gr 30, Gr 35 and Page (t)
Used German and foreign projectiles
Same as above
Used German and foreign projectiles
Used Czech design and named projectiles: 8.35 cm Gr 27/30 (t) and 8.35 cm Page (t)
Used British HE and smoke shells: Sprg Pan 100 and Sprg Pan 106 (a)
No information given
It fired various rockets used by the Navy, such as HE, flare, etc.
Used HE rockets, designated: 8.6 cm RSpgr L/45 and RSpgr L/35
Used British ammo: HE (Gr 292 & 295 (a)) and Smoke: Rauchgr (a)
Used ammo: HE (8.8 cm Sprg L/45, Sprg L/45, 282 & Sprg Pan L/45 K), AP (Page & Page 39), APC (Page Pan 84/2) and Inc-Shrapnel (Gr Br Schk Flak)
Used HE ammo: 8.8 cm Sprg L/45
Used HE and Star projectiles
Used HE and Star projectiles

References

9, p. 237
12
12
5a, p. 69
5, p. 69
5a, pp. 28-9
12
5a, pp. 70-9
5a, pp. 69 and 9, p. 69
5a, p. 70
5a, p. 70
9, p. 241
9, pp. 256-7
5a, p. 70
9, pp. 438, 444, 446 & 448
9, p. 444
5b, table 2
5b, table 2
5a, p. 41 and 9, pp. 444-5 & 448
9, pp. 438, 444, 446 & 448 and Ref 12
9, pp. 438, 444, 446 & 448
9, pp. 437-9, 441 & 444 and Ref 12
9, p. 441
12
5a, pp. 246-5, 9, pp. 442 & 447 and Ref 12

WEAPONS

(CALIBERS 88 mm TO 105 mm)



Ger 255 WEAPONS

[CALIBERS 128 mm to 300 mm]

128 mm RHEINMETALL
ANTITANK GUN
(128 cm K-44)

(15 cm sFH 18)
MOUNTED ON PzKw III
CHASSIS
(BUMBLE BEE)

150 mm MEDIUM
FIELD HOWITZER WITH
MODIFIED BREECH-LOADER
(15 cm sFH 18-43)

150 mm ASSAULT HOWITZER
(15 cm sFH 43) ON PzKw III
CHASSIS ("GRIZZLY BEAR")

150 mm MEDIUM INFANTRY
HOWITZER (15 cm sFH 33)

210 mm MORTAR (21 cm GrW 19)

210 mm GUN 3812 cm KANONE 381

240 mm GUN
MOD 3 (24 cm
KANONE 3)

150 mm SKODA MORTAR

280 mm RAILROAD GUN
NEPOMUCOPOLAND 33NNE
(28 cm KANONE 57E)

300 mm ROCKET LAUNCHER

150 mm MEDIUM FIELD HOWITZER
(15 cm sFH 18) WITH
MUZZLE BRAKE

Caliber and Designation

150 mm Gun 18: 15 cm K 18

150 mm Heavy Field Howitzers:
15 cm sFH 18, sFH 18/4, sFH 18/7
and sFH 36

150 mm Field Howitzer: 15 cm
sFH 40

150 mm Heavy Infantry Gun (Howitzer):
15 cm sFH 33 or sFH 33

150 mm Gun 39: 15 cm K 39

150 mm Railway Gun: 15 cm K (E)

150 mm Heavy Turbine Howitzer:
15 cm sHT

150 mm Heavy Field Howitzer:
15 cm sFH 42

150 mm Gun on Howitzer Carriage:
15 cm sFH 18/43 (with modified breech-
loader)

150 mm Heavy Field Howitzer:
15 cm sFH 18/43 (with modified breech-
loader)

150 mm Assault Howitzer:
15 cm sFH 43 (L/12)

150 mm Recoilless Gun:
15 cm LG 43

150 mm Caech Guns and Howitzers:
15 cm K 15/16 (i), sFH 14/160,
sFH 25 (i) and sFH 37 (i)

150 mm Rocket Launcher

152 mm (5.9") Rocket Launcher

152 mm Italian Heavy Field Howitzer:
15.2 cm sFH 412 (i)

152 mm Russian Guns and Howitzers:
15.2 cm sFH 404 (i), sFH 443 (i),
sFH 445 (i), KH 453/1 (i), KH 453/2 (i)
and KH 456 (i)

155 mm (6.10") Belgian Gun:
15.5 cm K 452 (b)

155 mm French Guns and Howitzers:
15.5 cm sFH 414 (i), sFH 415 (i),
K 416, 417, 418, 419, 420 and 425 (i)

155 mm Polish Heavy Field
Howitzer: 15.5 cm sFH 17 (p)

155 mm Yugoslav Guns and
Howitzers: 15.5 cm sFH 427/1
442/2 (i) and K 403 (i)

170 mm (6.69") Gun in Mortar
Mounting: 17 cm Kilmelaf

170 mm Gun 18: 17 cm K 18

170 mm Railway Gun: 17 cm K (E)

170 mm American Gun: 17 cm K (S)

172.6 mm (6.79") Naval Gun:
17 cm SF L/40

194 mm (7.64") French Railway Gun:
19.4 cm K 486 (E2)

Ger 256

(Weapons) (cont'd)

Remarks, Uses and Some Characteristics

Used ammo: HE (15 cm K 18 & 42),
ME-A/C (Gr 19 Rot Bel), and AP (PzSprg
L/3.7 mHb)

Used ammo: HE (15 cm KGr 18, Gr 19, &
Gr 36 FES), HE cast steel (Stggr 19),
HE-A/C (Gr 19B), Rocket Assisted (Gr 19),
Hoc (Gr 39 Hb), HE, Sabot (Sprg 42 TS), AP,
Sabot (Pzgr 39 TS) and Smoke (Gr 18Nb,
Gr 19Nb, Gr 19Nb & Gr 40Nb)

No description given

Used ammo: HE (15 cm Gr 19 & Jgr 38),
Roddied bomb (Snzgr 42) and Smoke (Jgr 38Nb)

Used ammo: HE (15 cm KGr 18, Sprg L/4.6
& KGr 42), A/C (Gr 19 RotBel), AP (Pzgr)
and SAP (HalsPzgr)

Used ammo: HE (15 cm KGr 18) and
AC (Gr 19 Ba)

Used ammo: HE (15 cm Gr 19 &
Gr 19 Sgr) and A/C (Gr 19 Ba)

Used same ammo as 15 cm sFH 18

Used ammo: HE (15 cm KGr 18, Sprg L/4.3,
Sprg L/4.6 & Sprg mHb), A/C (Gr 19
Rot Bel) and APC BC HE (PzSprg L/3.8 mHb)

Can be seen at the Museum of Aberdeen
Proving Ground, Md

Used ammo: HE (15 cm Jgr 38 FES) and
Hoc (Jgr 39 Hb/A)

No information given

Used Caech ammo

Used 15 cm HE, smoke and chemical rockets

Used HE rocket projectile

Used Italian HE ammo: 15.2 cm Sprg 412/11(i)

Used Russian design HE, Smoke and
Snpapool ammo

Used Belgian HE ammo: 15.5 cm Gr 420 & 426(b)

Used French ammo: HE and Hoc

Used Polish HE ammo: 15.5 cm Gr 14 & 15 (p)

Used Yugoslav HE ammo

Used ammo: HE (17 cm KGr 38 & 39),
Incendiary (BrGr 39), AP (Pzgr 43) and
Snp Shell (Leuchtschnecke)

Can be seen at the Museum of Aberdeen
Proving Ground, Md

Used HE ammo: 17 cm Sprg L/4.7

Same as above

Used HE, AP and Star projectiles

Used French HE, cast steel proj: 19.4 cm
Stggr 486 (i) and 487 (i)

References

- 5a, p 97 and
9, pp 486-7,
491 & 493
- 5a, pp 486-7,
9, pp 493-7,
9, 506-7 & 509
and Ref 12
- 5a, p 95
- 9, pp 486, 494-5,
497-8 & 502
and Ref 12
- 5a, p 98 and
9, pp 487, 493,
498 & 504-5
- 9, pp 493 & 498
- 5a, pp 95-6 and
9, pp 493, 500 & 507
- 5a, p 95
- 5a, pp 96-7
- 12
- 5a, pp 99; 9, pp
486 & 491 & Ref 12
- 5a, p 95
- 5a, pp 99-101 &
9, pp 485 & 489-90
- 9, pp 245-7
- 9, pp 247-8
- 5a, p 106
- 5a, pp 104-7 &
9, pp 510-12
- 5a, p 108
- 5a, pp 101-5
& 108
- 5a, p 101
- 5a, pp 107-8
- 5a, p 112 and
9, pp 516-17
- 12
- 5a, p 112
- 5a, p 112
- 5b, table T1
- 5a, p 113 and
9, p 517

(Weapons) (cont'd)

Caliber and Designation

200 mm (7.874") Light Spigot Mortar: 20 cm L/40 (20 cm leichter Ladungswerfer)
 200 mm Rocket Launcher
 203 mm (8.0") Railway Gun: 20.3 cm K (E)
 203 mm Russian Heavy Howitzers: 20.3 cm H 503 (t) & H 503/2 (t)
 203 mm Naval Gun: 20.3 cm SK C/34a
 209.3 mm (R.24") Naval Gun: 21 cm SK L/45
 210 mm (R.27") Gun: 21 cm K/12 and K 12 (E)
 210 mm Mortar: 21 cm Mrs 18 (Heavy Howitzer)
 210 mm Long Mortar: 21 cm Lg Mrs 18
 210 mm Mortar: 21 cm Mrs 19
 210 mm Gun: 21 cm K 38
 210 mm Gun: 21 cm K 39/40 & 39/41
 210 mm Gun: 21 cm K 42
 210 mm Krupp Gun: 21 cm K (Krupp)
 210 mm Rocket Launcher: 21 cm R/L 42 and others
 210 mm Czech Heavy Howitzer: 21 cm Mrs K/1
 210 mm (R.27") Gun, designated K 12 L/110 km range
 220 mm (8.66") French Gun: 22 cm K 132 (t)
 220 mm Norwegian Heavy Howitzer: 22 cm Mrs M/24 (t)
 220 mm Polish Gun: 22 cm Mrs (p)
 220 mm Yugoslav Howitzer: 22 cm Mrs (t)
 234 mm (9.21") Belgian Howitzer: 234 mm H 543 (h) 545/1 (h) & 545/1 (t)
 215 mm (9.37") Naval Gun: 24 cm SK L/40
 240 mm (9.44") Howitzer: 24 cm H 39
 240 mm Gun: Models 3 and 1b: 24 cm K 3 & K 1b
 240 mm (R.27") Gun (Railway): 24 cm L/40 (E)
 240 mm Naval and Seacoast Gun: 24 cm SK L/50
 240 mm Thunder Gun (Railway): 24 cm K 38 (E)
 240 mm Krupp Gun: 24 cm K L/46 (Krupp)
 240 mm Czech Gun: 24 cm K (t)

Remarks, Uses and Some Characteristics

Used HE and Smoke mortar bombs: 20 cm Vgr 40 and Vgr 40N
 Used 20 cm AA Rocket
 Used ammo: HE (20.3 cm Sprgr L/4.7), AP (Page L/7) and Base (L/40) (Gr)
 Used Russian A/C proj: 20.3 cm Gr 503/2 Be (t)
 Used HE, AP and Star projectiles
 Used HE and AP projectiles
 Used HE projectile: 21 cm Gr 35 and HE A/C (Gr 18 Be)
 Used A/C proj: 21 cm Gr 18 Be
 Used ammo: HE (21 cm Gr 17, 17ang, 18, 18 Sp) and HE A/C (Gr 18 Be)
 Can be seen at the Museum of Aberdeen Proving Ground, Md
 Used HE shell: 21 cm KGr 38
 Used ammo: HE (21 cm Gr 39 & 40), HE A/C (Gr 39 Be) and SAP (Half Page 39)
 No description given
 No description given
 Used for launching various rockets, such as 11 cm R/L, Vgr 42 Spr and R 1000 BS
 Used Czech ammo: HE (21 cm AGGr 35) and HE - High Capacity (MGr 35)
 Used HE projectiles
 Used French HE ammo: 22 cm Gr 534 (t) & 535 (t)
 No description given
 Used Polish ammo: HE (22 cm Gr 40) and SAP (Half Page)
 Used Yugoslav HE ammo: 22 cm Gr (t)
 No description given
 Used HE and AP projectiles
 Used HE and AP projectiles
 Used Czech HE (24 cm Gr 39 & 39 ang), SAP (Gr 39 Be or Half Page) and French case steel HE shell: Sgr 358/2 (t)
 Used HE shell: 24 cm Gr 35, Mod 3 gun. Can be seen at the Museum of Aberdeen Proving Ground, Md
 Used HE ammo: 24 cm Sprgr L/4.2 and L/4.5
 Used HE ammo: 24 cm Sprgr L/4.1 and L/4.2
 Same as above
 No description given
 Used Czech HE ammo: 24 cm Gr 25 (t) and Gr 40 (t)

References

5a, p 34 and 9, p 354
 5a, p 248
 5a, p 114 and 9, pp 20-2
 9, p 318
 5h, table 11
 5a, p 116
 5a, p 109; 9, p 322 & Ref 12
 5a, p 109 and 9, p 323
 12
 5a, pp 114-15 and Ref 12
 5a, pp 10-11
 5a, p 115
 5a, p 116
 9, pp 248-9, 255-6 & 259-60
 5a, p 117
 5h, table 12
 5a, p 117
 5a, p 118
 5a, p 119
 5a, pp 119-20
 5h, table 12
 5h, table 12
 5a, p 120
 5a, p 120 and Ref 12
 5a, p 121 and 9, pp 254-5
 5a, p 121
 5a, p 121
 5a, p 121
 5a, p 122
 5a, p 122 and 9, p 325

(Weapons) (cont'd)

Caliber and Designation

240 mm French Gun: 24 cm K (E) 557 (t) & K 558 (t)
 240 mm French Gun: 24 cm K 546 (t) & K 546 (t)
 240 mm Russian Howitzer: 24 cm H 544 (t)
 270 mm (10.9") French Coast Howitzer: 27 cm K/40 Mrs 585 (t)
 274 mm (10.76") French Railway Gun: 27.4 cm K (E) 591 (t) and K (E) 592 (t)
 280 mm (11.024") Howitzer: 28 cm H L/12
 280 mm Coast Howitzer: 28 cm K/40 H
 280 mm Short Barrel Gun (Railway): 28 cm K/40 (E)
 280 mm Long Barrel Gun (Railway): 28 cm Lg R/L (E)
 Note: According to Ref 5h, table 13, the short and the long barrel guns were 283 mm.
 280 mm Thunder Bomb Gun (Railway): 28 cm ThBk (E) or Bk (E)
 280 mm Gun, Model 3 (Railway): 28 cm K 38 (E), nicknamed "Lump" and "Angie Annie"
 280 mm Gun (Railway): 28 cm K 571 (E) and K 572 (E)
 280 mm Naval and Seacoast Gun: 28 cm SK L/50
 280 mm French Heavy Howitzer: 28 cm Mrs 601 (t) and 602 (t)
 280 mm Russian Howitzer: 28 cm H 545/3 (t) and H 607 (t)
 280 Rocket Launcher
 283 mm (11.142") Naval Gun: 28 cm SK C/26, C/34 & C/40
 300 mm (11.81") Self-Propelled French Mortar
 300 mm Rocket Launcher, New Type
 305 mm (12.00") Naval and Seacoast Gun: 30.5 cm SK L/50
 305 mm Czech Howitzer: 30.5 cm Mrs (t)
 305 mm Belgian Howitzer: (30.5 cm) 632 (b), 632 (t) & 632 (t) and Yugoslav Gun (M 638 (t))
 310 mm (12.597") Gun (Railway) Mount
 320 mm Rocket Launcher (No German designation is given)
 340 mm (13.387") French Gun: 34 cm K 675 (t)
 355 mm (13.977") Howitzer M-1
 355 cm M1, known also as M1 Gun. Note: According to Ref 5h, in M1, the M1 gun was 350 mm
 365 mm (14.37") Recoilless Gun: 36.5 cm G 104, developed during WW II by the Rheinwaffen-Diesel A-G
 370 mm (14.567") French Gun: 37 cm K 710 (t)

Remarks, Uses and Some Characteristics

Used French HE case steel shell: 24 cm Sgr 557 (t)
 No description given
 No description given
 No information available
 Used French HE ammo: 27.4 cm Gr 593, 594, 595 and 596 (t)
 Used HE shell: 28 cm Sprgr L/3.5
 Same as above
 Used ammo: HE (28 cm Sprgr L/4.1) and HEAP (P/Sprgr L/2.6)
 Used HE ammo: 28 cm Sprgr L/4.4
 Used HE, ammo: 28 cm Gr 59 mthgr 2
 Used ammo: HE (28 cm Gr 35 & Gr 42) and rocket-assisted (RGr L/4.7)
 Used HE ammo: 28 cm Gr 39/42 & Gr 42
 Used ammo: HE (28 cm Sprgr L/3.6) and AP (Page L/3.2)
 No description given
 No description given
 Used HE rocket proj: 28 cm WkRgr
 Used HE and AP projectile
 Can be seen at the Museum of Aberdeen Proving Ground, Md
 Used HE rocket proj: 30 cm Wk 42 Spr
 Used ammo: HE (30.5 cm Sprgr L/3.6) and AP (Page L/3.4)
 Used Czech ammo: HE (30.5 cm Gr 35) and HE - High Capacity (MGr 35)
 No description given
 Can be seen at the Museum of Aberdeen Proving Ground, Md
 Used HE rocket ammo designated as 32 cm Wk
 Used French ammo: HE, case steel (34 cm Sgr) and AP (Page)
 Used A/C ammo: 35.5 cm Gr Be, Rü (Rüchling) Gr 42 Be and RSGr 44 Be
 Note: According to Ref 5h, in M1, the M1 gun was 350 mm
 Not described here because Ref 8, v 3 is confidential
 No description given

References

5a, p 123
 5a, pp 123-5
 5a, p 122
 5a, p 124
 5a, p 124
 5a, p 124
 5a, p 124
 5a, p 125
 5a, p 125
 5a, pp 125-6
 5a, p 126 and 9, p 329
 5a, p 126; 9, pp 327-8 and Ref 12
 5a, p 127
 5a, p 127
 5a, p 128
 5a, p 127
 9, pp 248-51
 5h, tables 12 and 13
 12
 9, pp 251-3 and Ref 12
 5a, p 129
 5a, p 128
 5a, pp 129-30
 12
 9, pp 254-6
 5a, p 130
 5a, p 130; 9, p 328 & Ref 12
 8, v 3, pp 614 & 623
 5a, p 131

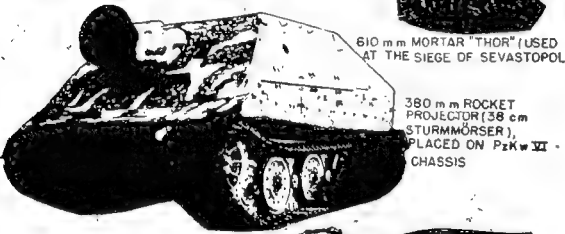
Ger 263
WEAPONS
(CALIBERS 310 mm TO 610 mm)



310 mm GLOTT GUN, MOUNTED ON 280 mm RAILWAY GUN [28 cm K 5 (E)] MOUNT



420 mm HOWITZER (WEIGHT OF WEAPON 35 TONS, WT OF PROJECTILE 1750 LB., RANGE 14500 YDS)
355 mm HOWITZER M1 BARREL (PLACED ON TRANSPORT CARRIER)

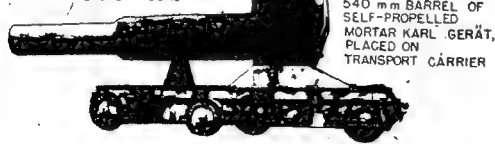


610 mm MORTAR "THOR" (USED AT THE SIEGE OF SEVASTOPOL)

380 mm ROCKET PROJECTOR (38 cm STURMMÖRSE), PLACED ON PRK M VII CHASSIS



540 mm SELF-PROPELLED MORTAR KARL GERÄT ON RAILROAD TRUCKS



540 mm BARREL OF SELF-PROPELLED MORTAR KARL GERÄT, PLACED ON TRANSPORT CARRIER

WP (Weißpulver) (Cultural or Prismatic Propellant). A black smokeless propellant in the form of small rectangular grains. It was first made under the name of WPC/89 (Weißpulver Construction 1899) by the Vereinigte Kün-
stliche Pulverfabrik in Rottweil, Württemberg for use in the Army gun, caliber 37 mm, 53 mm and 100 mm. The composition of WPC/89 was similar to the Italian Ballistone (Ref 1).

Bessert (Ref 2) gives the composition of an early WP as follows: NG 50, NC 50% and small quantity of DPBA, added.

Brennweit (Ref 3) gives for WP used after WW I: NG 38.5, NC 60, crocitolite or scarlate 1.0 and moisture 0.3%.

Reference: (1) J. Duval, Dictionnaire des Matières Explosives, Dunod, Paris (1902), p 811
(2) E. Bessert, Explosives, Van Nostrand, N.Y. (1919), p 253
(3) H. Brennweit, Das rauchlose Pulver, W. de Gruyter, Berlin (1926), p 156.

WPC/89. See under WP (Weißpulver).

W-Salt. The name given to Hexogen (RDX) prepd by the mixture of K methylenesulfonate (See under Hexogen).

Weißpulver. See WP.

Wingebühnen Geschütz. See Tapered Bore Gun.

X-4 was a first-stabilized guided missile with a proximity fused warhead developed especially for use by fighter planes against enemy bomber formations. It was propelled by a liquid fuel (Tosol 350) and an oxy-acetylene (Safel). Some experimental models were equipped with devices called "Kinnack" and "Pulack" [TM 5-1985 (1933), pp 215-19].

X-Ray Equipment (Röntgenrichtung). A short description of the x-ray equipment manufacturing industry is given in CIOS Report 28-31 (1943).

X-Rocket Guided Missiles. See Ruhrpawl under Guided Missiles.

X-Stoff. See Tetan.

Zebel, in 1899, constructed a metallic cartridge consisting of two compartments divided by a thin partition. In one of the compartments was a mixture of Ca carbide and Ba peroxide; while the other contained a dilute acid solution. On break the partition the acid reacted with carbide and peroxide to form a mixture of acetylene and oxygen which immediately exploded.

Reference: Duval, Dictionnaire (1902), p 814.

Zeitschur (Time Fuse), called in the U.S. Army Safety of Blasting Fuse. See under Fuses in the general section.

Zeitschurweitzünder (Time Igniter With Fuse). See under Electric Igniters, or Primers and also in Beyling-Dröckop (1936) pp 175 & 260-6.

Zell-ligant was a porous vinyl chloride polymer laminate. For use as an outside armor for the air intake tube (Schneidke) as well as for the peroxide in order to prevent the direction of submarines by short waves sent from enemy planes by radar.

The pores of Zell-ligant contained nitrogen generated within the material by a special process involving the use of a substance known as "Porofor N". For this a mixture consisting of polyvinyl chloride, 9% and Porofor N 5% was heated in an autoclave at 150° and then the mass was laminated. During this process the Porofor N dissolved in the vinyl chloride and reacted with the liberation of nitrogen which formed bubbles inside the material. Each Schneidke tube was covered with 7-8 layers of the above porous laminate, each layer being separated from the other by interposing carbon black coating paper, which was slightly conductive to electricity. It was assumed that the incoming short waves from a radar generated convection currents within the carbon paper and these currents were subsequently buffered if not completely absorbed by the laminae. Reflection of the short wave was thus minimized if not completely absorbed by the insulating mass.

Reference: CIOS Report 25-18 (1945), pp 29-30.

Zellpulch. See under Reusch's White Blasting Powder.

Zinn (Tin). See general section.

Noise According to A. Stettin, Spreng- und Schießstoffe, (1948), p 43, small quantities of tin, in oil, are easily reducible catalysts were incorporated in some German NG smokeless propellants in order to protect the interior of gun barrels from erosion.

Z-Stoff the name given to Na or Ca permanganate used as oxidizing components of rocket propellants in which it served as a combustible component. Z-Stoff was used in the Feuerlöcher type guided missiles called Hecht-Torpedos, Luchter etc N.Y. (1951), pp 45-46.

Z-Stoff C. An aqueous solution of calcium permanganate containing 600 g MnO₂ per liter 50 g LiCl at 20° and 10 g "Z". Used as a catalyst, as described below (CIOS 10-15, p 110).

Z-Stoff N. An aqueous solution of sodium permanganate containing 600 g of MnO₂ per liter 50 g LiCl at 20° and 10 g "Z". Used as a liquid catalyst in liquid rocket propellants to assist the decomposition of hydrogen peroxide which served as a source of oxygen (CIOS 10-15, pp 8 & 10).

Note: Z-Stoff N was used in summer since it is in "Z" while Z-Stoff C was used in winter (it is "Z"). When Z-Stoff C or N is used to decompose the Z-Stoff (hydrogen peroxide) the gaseous products contain besides water vapor and oxygen some small particles of manganese dioxide. Due to the presence of these particles, the gaseous mixture thus produced is not suitable for driving a turbine but can be used for other purposes such as in assisted take-off units and in missiles. When it is necessary to obtain a gaseous mixture free of MnO₂, the decomposition of H₂O₂ is conducted by means of a solid catalyst, such as described under WP 14.

Zünder. See Fuse.

Zündersprengkessel-43. A separate cap and detonator assembly designed for use in some A/T mines in conjunction with a tilt type igniter, called Kämpfer-43 [TM 5-1985-2 (1933)].

Zünderst. See Initiatorvermögen.

Zündsprühmisch. See Cartridge Case Percussion Primer.

Zündst. (Pinning Composition). See Primary and Initiating Compositions.

Zündschwurzünder (Igniter or Lighter for Fuse). Beyling-Dröckop (1936), pp 166-67, describes several types of igniters. Some of them are used for driving a turbine fire mines (für Schiffsversenkergruben), while others for gaseous mines (für Schiffsversenkergruben).

Zündstoffe oder Initiatorvermögen (Pinning, Igniter or Initiating Compounds). See Primer and Initiating Compositions.

Zündversenker (Ignition Intensifier). Ignition of a propellant in 50 to 280 mm weapon was accomplished by means of a primer combined with an igniter consisting of a black powder. For ignition, extremely called Zündversenker was fixed in front of the primer. This was filled with large grains of black powder and had a venturi in the forward end to blow the flame the full length of the charge. There were also one or two small side holes (igniter) near the charge as well.

Reference: CIOS 31-68 (1940), p 1. (See also under Igniters).

Zuschneider Drill. See Progressive Rilling.

Zusammengesetzte Zünder (Composite Igniter or Primers, as described in Beyling-Dröckop (1936), p 74).

Zwischenschladung, Zwischenladung, oder Zwischenzünder (Intermediate Charge or Booster) as described A. Stettin, Schieß- und Sprengstoffe, Leipzig (1933), p 334.

Zwischenzünder. See Zwischenladung.

Zwischenzündung. See Zwischenladung.

(In collaboration with H. A. Tisch and J. F. Neuck of
Picatinny Arsenal, Dover, New Jersey)

10

Mining(ore); dismantling(structure); abschliessen

[illegible][illegible]

[illegible]

Salary watch (lit Swedish)

[illegible]

Self-propelled (SP) mount;

- gun motor carriage (See also under Panzer)
- [Self-propelled gun]
- Subcaliber barrel for semi-automatic
- Semi-automatic rifle; self-loading rifle
- Semi-automatic pistol; self-loading pistol
- Semi-automatic weapon; self-loading weapon
- Semi-automatic rifle
- Self-destructing type of fuse
- Self-sustaining incendiary
- Spontaneous decomposition
- Spill
- Shipments, transshipment (Radioactive cargo) (CWS)
- vertical; perpendicular
- Sighting; lowering; hollow; sight
- Sighting; lowering; hollow; sight
- Production in series
- Permanent chemical warfare agent
- See Spitzgeschosse
- 'nadir; nadir
- Safety tank; draining tank
- Safety tank; draining tank
- Safety place; ammunition place
- Safety place; ammunition place
- Safety blasting powder
- Safety cylinder
- is made safe; lock (Out and Ammo)
- cores; protect; make safe
- See Spitzgeschosse
- Safety valve; safety valve
- Lock up
- Safety pin (Fz)
- Ammunition (Fz); end fire pin
- Safety fuse
- Fields; viability
- Field of view
- Screening; filter
- Boiling point
- 300 mm Railway Gun (See under Weapons)
- Signal
- Signal cartridge
- Signal rocket
- Ground signal projector
- Silence
- Deposited matter; sediment
- Security police
- See Schrapnellminen
- See Schrapnellminen
- Three-way adapter for 5-Mm
- Pedestal mount (G)

100

100

[illegible][illegible][illegible]

Poisted bullet with
 steel core; super
 Poisted bullet with
 core and thin trace
 Poisted bullet with
 and tracer
 Poisted bullet am
 Splint; cotter pin;
 Splinter; fragment
 Concrete fragment
 Fragment of shell
 (antipersonnel) bo
 Density of fragmen
 of shell fragments
 Fragmentation
 Fragmentation al
 over casting of the
 group of thin splinte
 Protective pagias
 splinterproof
 Trail speed (G) m
 Mph
 Megaphone
 Splint trail spade c
 Blasting job
 HE bomb; demolit
 HE bomb; demolit
 HE-HE high-we
 HE-HE bomb
 Demolition charge
 Demolition charge
 Demolition charge
 of a box containing
 Demolition service
 to blast
 Bombing (See al
 adumg)
 Fillety HE filling
 Explosive liquid
 Explosive liquid
 HE shell
 HE shell pattern d
 tapered nose gun
 HE shell in a curv
 (complete round of
 ammunition)
 See Sprengk
 Demolition (blast)
 Demolitor; blasting
 initiator
 Demolitor No. 8 (A
 Demolishing cord
 Demolition (blast)
 prepared demolition
 Prep demolition
 with delay 100 sec
 Prep demolition 10
 with delay 200 sec
 Demolition block;
 charge
 Demolition block;
 charge

LIST OF
GERMAN ABBREVIATIONS (Abkürzungen)
OF ORDNANCE AND RELATED TERMS

(in collaboration with K. F. Kempf at Aberdeen Proving Ground, Maryland)

▲

A-1	Abwehr	Défense
A-1f	Artillerie	Artillery
A/ben following projectile designation; white stencilling;	Ausammalsalung	Expelling charge of a shaped or smoke projectile
A/such as in: HL/A, HL/B, HL/C	Hohlladung A, B, and C	Types of hollow charges
A-1	Aggregat Eins	Aggregate No 1
Note: A-1 was the first successful liquid-propellant rocket developed at the Rocket Development Center at Kummersdorf West.	A-4	Aggregat No 4
Note: A-4, commonly known as V-2, was one of the most successful liquid-propellant rockets (See V-2 in the description section).	A-4	Aggregat No 4
a/a; AA	alte Art	Old type or pattern (See also aa and af)
AA	Abschussbehälter	Aerial bomb container
Examples: AB 23 SDZ; AB 24 SDZ; AB 36, AB 42, AB 500-B; AB 500-A, etc. [X 2789-2 (1953), pp 93-108 and 11-119]	Ausenschießer (Kassengragnate rot AB)	Smoke canister ejected from projectile on burst (Gun shell with red smoke canister)
AB (black stencilling on a projectile, such as EG rot AB)		
Abb	Abkürzung	Abbreviation
Abgr; Abg	Abgräber	Ricochet; ricochet burst
Abv	Abstristung	Demobilization; disarmament
Abw	Abweiser	Sender
abwbl	abschlen	shoot down
Abzsch Ger	Abzschungsgerät	Grenade launcher
ABSt	Artillerie Beobachtungsstelle	Artillery Observation post
Abt; Abtg	Abteilung	Section; detachment; department
Abz	Abwehr	Défense
ac	achse	Trigger; trigger of the current year
ACB (such as in WGR at ACB)	achse currentis	Marking on a plastic PDF/A in 80 mm smoke mortar shell [TM 9-1985-3 (1953), p 591]
WGB at ACB	(Waffenarsenalmeister Treiblad ACB)	on bar, at the twisted
w/g	an der	Through official sources; through channels
aD	nasser Dienst	Army dagger
aDD	an dem Dienstwege	Adjutant
AD	Annehmlich	Admiral
Adj	Adjutant	General Service Regulations
Adm	Administ	Address
ADO	Allgemeine Dienstordnung	Ether
Adresse	Adresse	
Aether; Äther	Aether; Äther	
AEG	See under Vanplunius (descriptive section)	
ag; av	ausserer (äusser) Weize	distant diameter
AG	Artilleriesflieger	Artillery air observer; Artillery spotting filter
A/G	Atomgewicht	Atomic weight
A-G	Attingersalltschaft	Joint Stock Company; Open Corporation
AGFA; Agfa	A-C für Auslieferungsbezeichnung	Autoline Dye Manufacturing Corporation
AGF	Ausschussgeschosse	Sighting projectile
ANH	Allgemeines Heeresamt	-General Army Office
ANA	Armeehauptquartier	Army Headquarters
AK	Arbeitskörper	Army Corp
akt	aktiv	active; on duty
Al	Aluminium	Aluminum
Al (black stencilling following the designation of shell 7.5 cm GebGr 15 AU)	Aluminiumgranate	Designation of an HE shell containing some granular Al flash producer

ALF	See Under Wasplasma, etc (descriptive section)	Alcohol; ethyl alcohol; ethanol
Am	America; amerikaisch	America; American (See also VSUA)
Am	Ammonalpetre	Ammonium nitrate
Ammonatp	am Main	Am nitrate strip propellant on the Main (river)
a/M	am dem	at; by; on; near to
AML	Armer-Mundstueckelager	Army Ammunition Depot
AmSt	Amperestunde	Amper-hour
amsl	amlich	official
As	Anisoi	Troismitrailleuse (TNAs)
As 60/40	Anisoi 60/40	TNAs 60 and Am nitrate 40%
Asl	Anfang	Beginning
AsfGeschw	Anfangsgeschwindigkeit	Initial velocity; muzzle velocity
Angew Chem	Angewandte Chemie (formerly Zeitschrift für Angewandte Chemie)	Applied Chemistry (Journal)
Ash	Anhang	Appendix; supplement
Asht	Anhänger	Trailer; supporter; follower
AshtV	Anhangswagen	Trailer
Asl	Anhalt	Plant; establishment
Asm; Asmerit	Anaerobung	Respirator; locomotor
Asn	Annahme	Acceptance; receipt
Asn	Ansehen der Chemie	Annals of Chemistry (Journal)
Asp	Anpassung	Adaptation
ANR	Armes Nachrichten Regiment	Army Signal Regiment
asoch	anschießen	to hit by shooting
Ansch Puz	Anschiespatrone	Ammo used for adjustment fire
Asst	Ansaat	Establishment; institution
Asa	Ansaat	Number
Asa	Ansaat	Indicator; informer
ANZ	Ansaender	Igniter
ANZ-29	Ansaender 29	Friction pull type igniter used to ignite a safety fuse or to set off a smoke candle (TM 9-1983-2, pp 286-7)
	Artillerieoffizier	Artillery officer
AO	Agro	Designation of airplanes assigned by Ago Co
As	Artilleriepunkt	Artillery reference point (gunnery)
App	Apparat	Apparatus; device; equipment
AR	Artillerieregiment	Artillery regiment
As AR	Arado	Designation of airplanes assigned by Arado Co
ARDA	Meaning unknown to us	Designation of a smoke signal flask (TM 9-1983-2, p 80, Fig 84)
A/R	am Rhein	on the Rhine (river)
AsR	Arbeitskummission	Ammo for use in Arctic climate
Arm	Armo	Army (formation above Army Corps)
arm	armiert	Armed
Arm; As	Armenal	Armenal
Arm	Arrenal	Armenic (As)
Arm; Art; A	Artillerie	Artillery
AS SCMP1	Artilleriegeschwaderplatz	Artillery firing range; Proving ground
AS	Anforderungssignal	Call signal
ASG	Anforderungsstelle	Computing station (sound and flash ranging gunnery)
At; Am	Atmosphäre	Atmosphere
Att	Anschlag	Attachment
Atu	Attrappe	Dummy
Atu	Atmosphärenüberdruck	Gage pressure; pressure above atmospheric
Atube	Atmosphärensensor	Overhead sensor
Atub	Aufbau	Building up; construction; organization
AtuG	Aufilage	Edition
Atub	Aufnahme	Photographic picture
Atub	Aufschiff	Up set (gunnery)
Atub	Aufschuss	Yield
Atub	Aufbildung	Training
Atub	Aufbrechung	Erosion (of a barrel)
Atub	Aufführung	Execution; completion; model; design
Atub	Ausgabe	Issue; issuance
Atub	Auslastung	Arm and equipment

[illegible]

Ventral gun mount
Light case shell of cast steel (TM 9-1985-3, p 349)
Fixed round with a smoke producing projectile used
for adjustment fire
Fire: incendiary

See 943

22. **Examiner** Pull type lighter
Examiner 22-35 (pull type lighter used with help since in opening various places) and 22-43 (pull or pressure type lighter, used in bulky maps and small pieces) (TSP-1949-2, pp. 289 & 293)
or
22a, 22b. **See 249** at the examiner's table
22c. **Zinc-oxide-chloride** Fine granular cap
22d. **Zinc-oxide** Cylindrical powder
22e. **Diamonds** Diamonds

Abbreviations See at the end of the previous section: "Vocabulary of Common Abbreviations, Acronyms and Related Terms"
References